


Article

Anti-Inflammatory Diet and Protein-Enriched Diet Can Reduce the Risk of Cognitive Impairment among Older Adults: A Nationwide Cross-Sectional Research

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Abstract: Background: Cognitive impairment (CI) is a common mental health disorder among older adults, and dietary patterns have an impact on cognitive function. However, no systematic researches have constructed anti-inflammatory diet (AID) and protein-enriched diet (PED) to explore their association with CI among older adults in China. Methods: The data used in this study were obtained from the 2018 waves of the China Longitudinal Health and Longevity Survey (CLHLS). We construct AID, PED, and calculate scores for CI. We use binary logistic regression to explore the relationship between them, and use restrictive cubic splines to determine whether the relationships are non-linear. Subgroup analysis and sensitivity analysis were used to demonstrate the robustness of the results. Results: A total of 8692 participants (mean age is 83.53 years) were included in the analysis. We found that participants with a higher AID (OR = 0.789, 95% confidence interval: 0.740–0.842, $p < 0.001$) and PED (OR = 0.910, 95% confidence interval: 0.866–0.956, $p < 0.001$) score showed lower odds of suffering from CI. Besides, the relationship between the two dietary patterns and CI is linear, and the results of subgroup analysis and sensitivity analysis are also significant. Conclusion: Higher intakes of AID and PED are associated with a lower risk of CI among older adults, which has important implications for future prevention and control of CI from a dietary and nutritional perspective.

Keywords: anti-inflammatory diet; cognitive impairment; older adults; protein-enriched diet



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1. Introduction

With the rapid growth of China's aging population and longer life expectancy, age-related disorders have become a major public health problem [1]. One of the worrying issues is the deterioration of cognitive function (CF) [2], which may not only seriously harm individuals' well-being, but also impose a huge burden on their families and caregivers, as well as the financial and health systems of the society [3]. CF declines with age [4], age-related cognitive impairment (CI) is a common disorder in older adults, ranging from mild CI to severe dementia [5], and with the aging of the global population, dementia is an increasingly serious public health issue [6]. It is expected that the number of dementia patients will reach 82 million by 2030 and 152 million by 2050 as the population ages [7]. The cost of caring for dementia patients worldwide exceeds USD 800 billion annually and is expected to increase to USD 2 trillion by 2030 [8]. China has the largest number of dementia patients in the world (9.5 million), followed by the United States (4.2 million) [9]. According to a nationwide survey conducted in China from 2015 to 2018, approximately 15.5% of the population aged 60 and above suffered from mild CI, 6.0% suffered from dementia, and

3.9% had Alzheimer's disease [10]. Due to the incurable nature of dementia, it is crucial to detect CI in the early stages, and maintaining CF is a core component of successful aging [11].

The Lancet Commission concluded that lifestyle factors, including diet and nutrition, could prevent or delay 40% of dementia worldwide [12]. For instance, in terms of physical activity, a study has found that an increase in exercise and resistance exercises are beneficial for CF [13], and they can reduce the risk for dementia [14]. What is more, a combination of aerobic and strength training can slow the decline in motor and cognitive abilities in people with dementia [15]. In the aspects of nutrition and diet, a study proved that beta-carotene is a powerful antioxidant and dietary precursor of vitamin A, playing an important role in maintaining mental health and CF [16]. Moreover, dietary polyphenols have a neuro-protective effect and may prevent or improve CI [17], and there is emerging evidence to suggest that several plant-based foods rich in polyphenols have a promoting effect on brain health [18]. Omega-3 fatty acids, found in some foods such as fish, legumes, and nuts, also have an impact on CI [19]; Omega-3 fatty acids include eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA) [20], which are an important factor affecting the health of the brain structure and function [21]. Therefore, implementing a healthy eating pattern is a potential strategy for preventing CI, which can alleviate CI that occurs with age [22]. As is well known, there is a certain relationship between some nutrients in the diet and inflammation [23]. The Dietary Inflammatory Index (DII) is a comprehensive indicator calculated based on the correlation between nutrients and systemic pro-inflammatory cytokine levels, which has the function of evaluating the likelihood of dietary inflammation [24]. Higher DII scores illustrate more pro-inflammatory diets, while more negative values indicate more anti-inflammatory diets [25]. The characteristic of aging is an increase in the concentration of many pro-inflammatory molecules in the circulation, a phenomenon known as "inflammation" [26,27]. Systemic inflammation can lead to cognitive decline and dementia by inducing a reactive pro-inflammatory environment in the central nervous system [28], and research has proved that higher pro-inflammatory dietary potential is associated with increased incidence of dementia [29]. Simultaneously, an anti-inflammatory diet (AID) may have an impact on some mental health disorders. Previous studies have found that AIDs can serve as potential interventions for depression [30], and adopting the interventions of anti-inflammatory and diversified diets may alleviate the burden of depression among older adults [31].

Earlier studies have found that many dietary patterns are associated with improved CF. A *JAMA* article showed that adding olive oil or nuts to the Mediterranean diet was associated with improvements in CF in older adults [32], and a number of other related studies have further confirmed the significant association between the Mediterranean diet and CF [33–35]. Additionally, the Mediterranean-DASH Intervention for Neurodegenerative Delay, known as the MIND diet, is a hybrid of the Mediterranean diet and the DASH (Dietary Approaches to Stop Hypertension) diet, which can prevent and slow the cognitive decline in older persons [36–39]. A prospective cohort study found that AID patterns are associated with decreased grip strength [40], and research has found that grip strength is associated with cognition and dementia [41,42], so there may be a certain connection between AID and CI. In the elderly population in northern China, a pro-inflammatory diet is associated with an increased risk of mild CI [43]. Despite numerous studies on other dietary patterns and CI, as well as studies on AID and other outcomes variables, research on the relationship between AID and CI among older Chinese adults remains undefined, which requires further research and certification.

Protein is a critical nutrient for normal CF [44], as some research suggested that protein intake is associated with better CF performance [45,46]. Creatine is found in many protein-rich meats, such as beef, lamb, chicken, and fish, and studies have found that creatine has a positive effect on brain health [47] and cognitive performance [48]. This may also be one of the reasons why protein-rich diets have a positive effect on CF. A balanced protein diet is beneficial for the CF of older adults in Japan [49]. Besides, a study has found

that the dietary protein intake, total animal protein intake, total meat, egg, and legume protein intake of adults aged 60 and above in foreign countries are positively correlated with CF [50]. Studies have shown that a protein-enriched Mediterranean diet fights malnutrition and promotes healthy neurocognitive aging in older adults [51]. Moreover, fish products are recommended as a protein dietary source and are associated with a lower risk of CI [52], while milk and dairy intake in middle age may have a protective effect against CI [53]. Nonetheless, other studies reported that there is no significant correlation between protein intake and CF [54,55]. Considering that previous research on the relationship between protein intake and CF were not entirely consistent, and no one had built a systematic protein-enriched diet (PED) to explore its relationship with CI, we further explored the association between them among older adults in China by using a systematic dietary pattern.

Based on the analysis above, since the role of protein and anti-inflammatory diets on CI is unclear, there is an increasing need to construct two dietary indices for measuring their relationship with CI. Therefore, in this study, we will construct two dietary indices, AID and PED, and explore their effects on CI among Chinese older adults based on the CLHLS database.

2. Materials and Methods

2.1. Participants and Process

The data used in this study were obtained from the 2018 waves of the China Longitudinal Health and Longevity Survey (CLHLS), which used a multi-stage stratified cluster sampling design in 23 out of 31 provinces in China. In this research, we used the data of the 2018 wave to investigate the association between AID, PED, and CI. We excluded 503 cases of missing dietary index data, 417 cases of missing CI data, as well as 4560 individuals with missing covariate data. After eliminating the merged missing values, the final analysis sample utilized in this study contained 8692 adults aged 60 years or older. The specific data cleaning process is shown in Figure 1. All subjects signed informed consent for baseline and follow-up surveys. The project was approved by the Biomedical Ethics Committee of Peking University, China (IRB00001052-13074).

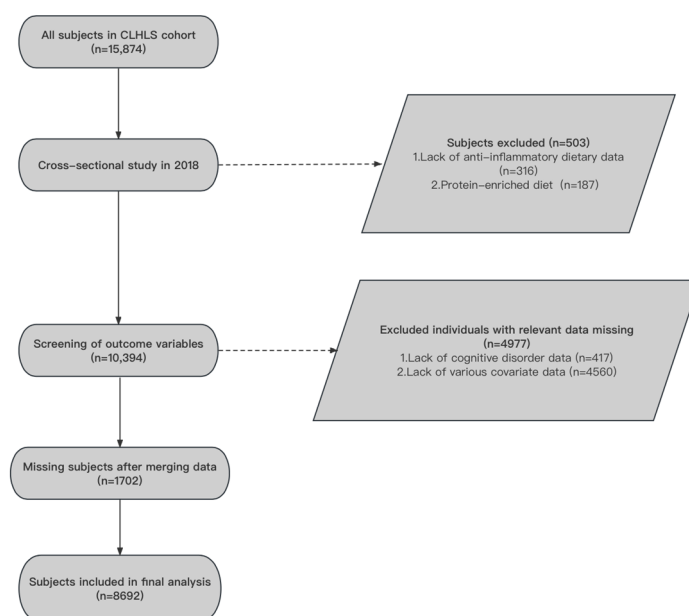


Figure 1. Data cleaning flow chart.

2.2. Assessment of Dietary Index

Five protein-enrich food sources were considered in this research, which are meats, fish, eggs, dairy and its products, and bean products. We constructed the PED, which

ranged from 0 to 5, by summing up the frequency of intake of protein-enriched food. The details of protein-enriched foods have been described in an earlier study [24]. What is more, the anti-inflammatory food included vegetables, fruits, legumes and their products, nuts, and tea, and they were used to establish the AID, ranging from 0 to 5. The consumption of any food category above “frequently or virtually every day” was deemed to equal one AID unit [31], and the AID scores are obtained by summing the scores of each food consumed.

2.3. Assessment of Cognitive Impairment

The CI of CLHLS participants was assessed by the Chinese version of the Mini-Mental State Examination (MMSE) through a home-based interview, which includes 24 items, covering 7 subscales including orientation (4 points for time orientation and 1 point for location orientation); naming foods (naming as many kinds of food as possible in 1 min, 7 points); registration of 3 words (3 points); attention and calculation (mentally subtracting 3 iteratively from 20, 5 points); copying a figure (1 point); recall (delayed recall of the 3 words mentioned above, 3 points); and language (2 points for naming objectives, 1 point for repeating a sentence, and 3 points for listening and following directions). The MMSE score ranges from 0 to 30, and higher scores represent a better CF. The validity and reliability of this Chinese MMSE have been validated in previous studies [56,57]. The judgment of CI is related to the level of education received: illiteracy scores ≤ 17 , primary school level scores ≤ 20 , and high school or above scores ≤ 24 are considered to have CI.

2.4. Covariates

In order to minimize their impact, we controlled for a large number of potential confounding factors, including age, gender (male or female), residence (urban or rural), nationality (“Han” or “other”), and education level (0 year, 1–6 years, and ≥ 7 years). Smoking and alcohol consumption were binary variables (yes or no). The calculation method for body mass index (BMI) was to divide weight (kilograms) by the square of height (meters). In addition, we also classified marital status into four categories: unmarried, married, divorced or separated, and widowed, and assigned scores of 0, 1, 2, 3. Assigned a score of 0 to “good”, “very good”, and “average” in the self-assessment of life satisfaction and health condition, and assigned a score of 1 to “not good” and “very bad”. Finally, we categorized sleep duration into three levels: less than 7 h, 7–9 h, and more than 9 h, with scores of 0, 1, 2, respectively.

2.5. Statistical Analysis

Continuous variables that follow a normal distribution are represented by mean \pm standard deviation (SD). Categorical variables are represented in terms of frequency and percentage. Analysis of variance or Chi-square test is used to compare differences among subjects under different demographic characteristics. We use a binary logistic regression model to evaluate the association of CI with PED and AID separately. Four hierarchical regression models have been established: the basic model (Model 1) does not include any covariate; Model 2 controlled age, gender, health condition, life satisfaction, and sleep duration; Model 3 further controlled for smoking, drinking, and marital status; Model 4 additionally controlled BMI, nationality, and residence.

Restricted cubic splines (RCS) are adept at handling the non-linear relationship between continuous variables and response variables. We used RCS to determine whether there is a non-linear relationship between AID, PED, and CI. Additionally, we performed subgroup analysis for the decline of CF, grouped by gender, residence, sleep duration, health condition, and life satisfaction, and tested for interactions between grouping variables and two dietary patterns using likelihood ratio tests. We also adopted sensitivity analyses using the full model (Model 4) under different participants. All statistical analyses were completed using SPSS 26.0 and R 4.3.0. $p < 0.05$ indicates statistical significance in this study.

3. Results

3.1. The Characteristics of Study Participants

In this study, we used information from 8692 participants, whose mean age was 83.53 ± 11.48 . The gender ratio was relatively balanced (44.8% male and 55.2% female), and their nationality was mostly Han (95.1%). More than half of the population had a BMI value between 18.5 and 23.9 (54.5%). In addition, the vast majority of them lived in rural areas (82.6%) and had a relatively low level of education, with only 20.6% receiving more than six years of education. Among them, 15.9% smoked, 14.9% drank alcohol, and more than half of the older adults slept less than 7 h (53.2%). They had a good self-evaluation of life satisfaction and health condition, accounting for 97% and 86.5%, respectively. The AID, PED, and CI scores were significantly different among these subgroups: BMI, gender, education, marital status, drinking, sleep duration, and health condition. More detailed information can be found in Table 1.

Table 1. Basic demographic characteristics of the participants.

Variable	Frequency (%)	AID (M \pm SD)	p Value	PED (M \pm SD)	p Value	CI (M \pm SD)	p Value
Age, years	8692 (100%)	2.32 ± 1.18	$p < 0.001$	2.95 ± 1.44	0.093	25.28 ± 6.01	$p < 0.001$
BMI, kg/m ²							
<18.5	1304 (15%)	2.05 ± 1.12	$p < 0.001$	2.82 ± 1.38	$p < 0.001$	22.65 ± 7.42	$p < 0.001$
18.5–23.9	4740 (54.5%)	2.27 ± 1.18		2.91 ± 1.45		25.25 ± 5.97	
23.9–27.9	2145 (24.7%)	2.54 ± 1.18		3.11 ± 1.43		26.68 ± 4.80	
>27.9	503 (5.8%)	2.49 ± 1.15		3.01 ± 1.45		26.46 ± 4.57	
Gender							
Male	3896 (44.8%)	2.47 ± 1.22	$p < 0.001$	3.08 ± 1.39	$p < 0.001$	26.54 ± 4.97	$p < 0.001$
Female	4796 (55.2%)	2.19 ± 1.14		2.84 ± 1.46		24.27 ± 6.57	
Nationality							
Han	8262 (95.1%)	2.33 ± 1.19	$p < 0.001$	2.98 ± 1.44	0.024	25.30 ± 5.99	0.066
Others	430 (4.9%)	2.03 ± 1.08		2.45 ± 1.27		24.92 ± 6.52	
Education (years)							
0	3940 (45.3%)	1.98 ± 1.06	$p < 0.001$	2.65 ± 1.43	$p < 0.001$	22.78 ± 6.87	$p < 0.001$
1–6	2965 (34.1%)	2.34 ± 1.14		2.94 ± 1.38		26.94 ± 4.37	
>6	1787 (20.6%)	3.03 ± 1.18		3.63 ± 1.31		28.07 ± 3.73	
Marital status							
Unmarried	68 (0.8%)	1.93 ± 1.14	$p < 0.001$	2.46 ± 1.49	$p < 0.001$	25.32 ± 5.18	$p < 0.001$
Married	3854 (44.3%)	2.51 ± 1.21		3.05 ± 1.42		27.53 ± 3.74	
Divorce or Separation	184 (2.1%)	2.47 ± 1.21		3.17 ± 1.41		26.88 ± 4.50	
Widowed	4586 (52.8%)	2.16 ± 1.13		2.87 ± 1.45		23.33 ± 6.89	
Smoking							
Yes	1379 (15.9%)	2.40 ± 1.19	0.275	2.94 ± 1.38	0.017	26.60 ± 4.83	$p < 0.001$
No	7313 (84.1%)	2.30 ± 1.18		2.95 ± 1.45		25.04 ± 6.18	
Drinking							
Yes	1292 (14.9%)	2.56 ± 1.23	$p < 0.001$	3.11 ± 1.37	0.007	26.58 ± 4.81	$p < 0.001$
No	7400 (85.1%)	2.28 ± 1.17		2.92 ± 1.45		25.06 ± 6.17	
Sleep Duration (h)							
<7 h	4628 (53.2%)	2.31 ± 1.20	$p < 0.001$	2.91 ± 1.48	0.014	25.71 ± 5.58	$p < 0.001$
7–9 h	2451 (28.2%)	2.40 ± 1.16		3.02 ± 1.40		26.04 ± 5.40	
>9 h	1613 (18.6%)	2.22 ± 1.14		2.97 ± 1.34		22.91 ± 7.36	
Life Satisfaction							
Good	8431 (97%)	2.34 ± 1.18	0.096	2.98 ± 1.43	0.202	25.39 ± 5.93	$p < 0.001$
Not Good	261 (3%)	1.68 ± 1.10		2.19 ± 1.50		22.01 ± 7.60	
Residence							
Urban	1512 (17.4%)	2.95 ± 1.19	0.485	3.71 ± 1.31	$p < 0.001$	26.58 ± 5.37	$p < 0.001$
Rural	7180 (82.6%)	2.19 ± 1.14		2.79 ± 1.41		25.01 ± 6.11	
Health Condition							
Good	7516 (86.5%)	2.37 ± 1.18	$p < 0.001$	3.00 ± 1.42	$p < 0.001$	25.52 ± 5.83	$p < 0.001$
Not Good	1176 (13.5%)	2.00 ± 1.16		2.67 ± 1.50		23.78 ± 6.90	

Notes: AID: anti-inflammatory diet, PED: protein-enriched diet, CI: cognitive impairment, SD: standard deviation.

3.2. Association between AID, PED, and CI

In this study, we found that among older adults in China, the higher the AID (OR = 0.694, 95% confidence interval: 0.657–0.734, $p < 0.001$) and PED (OR = 0.902, 95% confidence interval: 0.864–0.941, $p < 0.001$) dietary scores, the lower the odds of suffering from CI in Model 1 without controlling any covariates. After controlling some demographic

variables (age, gender, health condition, life satisfaction, sleep duration, smoking, drinking, marital status, BMI, nationality, and residence) hierarchically in Model 2 to Model 4, the association was reduced, but remained statistically significant. Moreover, in the full model (Model 4), taking into account all the relevant covariates, AID (OR = 0.789, 95% confidence interval: 0.740–0.842, $p < 0.001$) and PED (OR = 0.910, 95% confidence interval: 0.866–0.956, $p < 0.001$) were still related with a reduced risk of suffering from CI, by 21% and 9%, respectively. More detailed information is provided in Table 2.

Table 2. Associations of AID, PED with CI among Chinese older adults.

Model	AID	PED
Model 1	0.694 (0.657, 0.734) ***	0.902 (0.864, 0.941) ***
Model 2	0.795 (0.746, 0.846) ***	0.914 (0.871, 0.960) ***
Model 3	0.798 (0.749, 0.850) ***	0.917 (0.874, 0.962) ***
Model 4	0.789 (0.740, 0.842) ***	0.910 (0.866, 0.956) ***

Notes: AID: anti-inflammatory diet, PED: protein-enriched diet, *** $p < 0.001$.

3.3. Restricted Cubic Splines in the Regression Model

RCS did not reveal any significant non-linear relationship between two dietary patterns and CI in the full model (AID: $p_{\text{overall}} < 0.001$, $p_{\text{non-linear}} = 0.063$; PED: $p_{\text{overall}} = 0.001$, $p_{\text{non-linear}} = 0.421$). Figure 2 shows the risk of CI decreased when AID or PED increased. We can see that in Part A, the risk of CI decreases rapidly with the increase in the AID scores before the OR value drops to 1. As can be seen from Part B, the CI risk also shows a decreasing trend with the increase in PED scores, but its overall trend is relatively gentle compared to AID. Furthermore, we found that the protective effect on CI was significantly increased when the intake dose of both diets exceeded 3 units, namely that it is recommended to eat more than three types of protein-enrich foods and anti-inflammatory foods regularly every day to prevent CI.

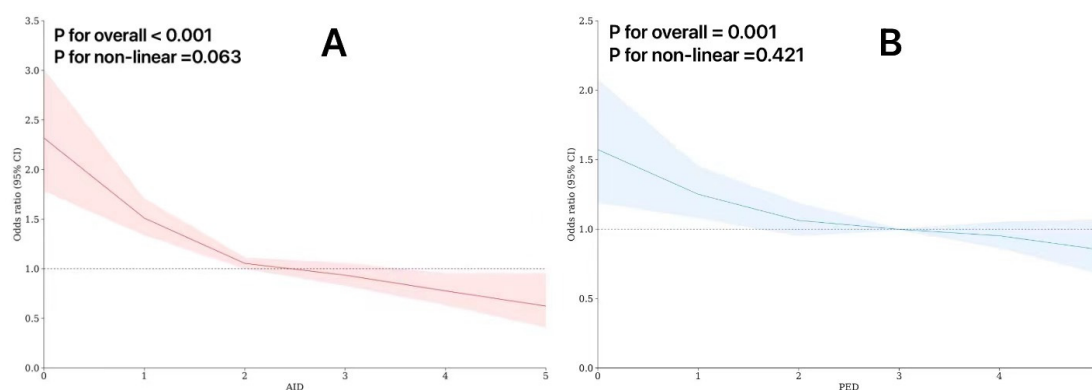


Figure 2. Restricted cubic spline for testing the hypothesis of non-linear correlation between AID, PED, and CI. (A) The linear relationship between AID and CI; (B) the linear relationship between PED and CI.

3.4. Subgroup Analysis

Figures 3 and 4 present the results of subgroup analysis. Based on subgroup analysis, we found that these associations are robust in subgroups of CI risk variables such as gender, residence, sleep duration, health condition, and life satisfaction. In addition, statistically significant interactions between residence, sleep duration, health condition, life satisfaction, and two dietary patterns (AID, and PED) on CI were observed (p -Interaction < 0.01). We can see from the figure that both dietary patterns still have a protective effect on CI under different subgroups, with all OR values below 1 and $p < 0.001$.

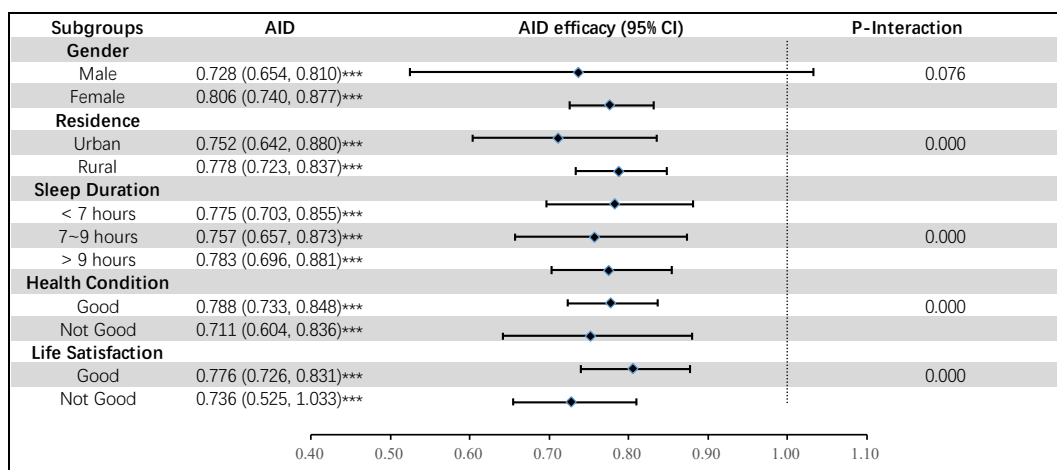


Figure 3. Associations of AID with CI among subpopulations. Notes: AID: anti-inflammatory diet, *** $p < 0.001$.

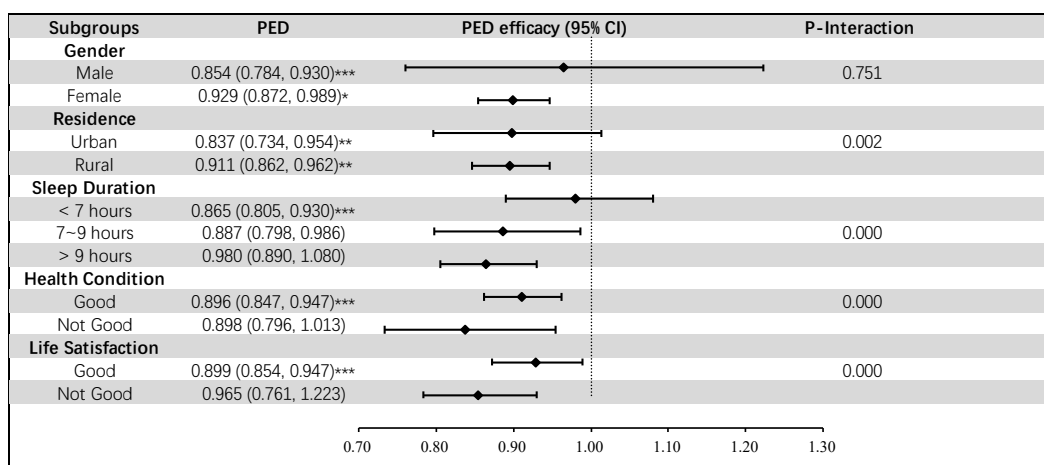


Figure 4. Associations of PED with CI among subpopulations. Notes: PED: protein-enriched diet, *** $p < 0.001$, ** $p < 0.01$, * $p < 0.05$.

3.5. Sensitivity Analysis

We performed sensitivity analysis using the full model (Model 4) under different participants. All of the associations between AID, PED, and CI are significant and consistent with the main results in this research. Exclusion criteria for sensitivity analysis: (1) excluded the older adults who sleep less than 7 h every day; (2) excluded the participants with poor health condition.

4. Discussion

The findings of this study are based on a nationwide population of older adults in China. We found that older adults with higher AID and PED scores are less likely to suffer from CI. Furthermore, we used RCS to determine whether there was a non-linear relationship between these two dietary patterns and CI, and demonstrated the robustness and reliability of our research results through subgroup analysis and sensitivity analysis.

Previous research has confirmed a number of risk factors that can impair normal CF, such as advanced age, lower levels of education, and unhealthy lifestyles, as well as physical and mental health disorders [58]. Our results are consistent with previous studies; people who have received education for a longer period of time have higher MMSE scores, which also confirms they have better CF due to the longer education. We found that a difference in CI exists among females and males ($p < 0.001$), similar to many previous studies [59,60].

This may be due to the previous differences in socio-economic status between men and women in society, which resulted in fewer opportunities for women to receive education, leading to older women becoming a more vulnerable group to CI [57]. In this study, there is also a significant gender difference in the scores of the two dietary patterns; studies have proved significant differences in dietary intake between men and women [61]. Therefore, one potential reason for this difference in the relationship between dietary patterns and CF may be gender [62]. Based on our exploration of the impact of dietary patterns on CI, it is reasonable that gender differences in dietary intake can lead to gender differences in CI. Moreover, we found a significant difference in CI scores with smoking, which is consistent with previous studies; current smoking may be positively correlated with CI in middle-aged people [63]. In older women, habitual sleep duration predicted future risk of CI, including dementia, independent of vascular risk factors [64], and individuals with insufficient sleep time (≤ 4 h per night) or excessive sleep time (≥ 10 h per night) may have an impact on CF [65,66]. Those explain why there were significant differences in CI by gender and sleep duration in this study. An Indian study found that mental health disorders such as CI and depressive symptoms in older adults differed in rural/urban areas [67], perhaps due to differences in socio-economic and educational levels between rural and urban areas. Additionally, our research findings have indicated that married older adults had significantly higher CF scores than other groups. Marriage is associated with a lower likelihood of CI; it provides social support, companionship, and participation in mental stimulation activities, which can help improve cognitive health [68]. Conversely, divorced and widowed older adults are particularly susceptible to CI [69]. In this study, individuals with better self-evaluation of life satisfaction and health condition had better CF. Satisfaction with life may be a positive psychological resource for maintaining CF and preventing the risk of dementia [70]. Some studies have found that sub-health condition may be a risk factor for CI in northern Chinese population. Early screening of sub-healthy individuals, as well as emergency treatment of sub-healthy individuals, may contribute to the prevention of CI [71].

Our research findings support evidence from a previous study on older adults that nutrition can alter the risk of future CI and dementia [72]. Some inconclusive evidence exists (mainly from observational studies and rarely from clinical trials) indicating a protective association between certain nutrients (such as folate, flavonoids, vitamin D, and certain lipids) or food groups (such as seafood, vegetables, and fruits, as well as potential moderate alcohol and caffeine intake) and cognitive outcomes in older adults. Various elements of the diet may be linked to CI and dementia; research has shown that a higher intake of monounsaturated fats can prevent mild CI in males and females aged 60–64 [73], and in people aged 50 years, intake of polyunsaturated fats was associated with a reduced risk of dementia [74]. A meta-analysis found a negative dose–response relationship between serum vitamin D concentration and the risk of developing dementia or Alzheimer’s disease [75]. There are studies reporting that consuming fish has an overall protective effect on cognitive decline [76]; this may be because fish products are rich in creatine. Another study found that among participants aged 60 and above, those who frequently consume milk and dairy products have a reduced risk of developing Alzheimer’s disease [77]. Fish, milk, and dairy products are all protein-enrich, which indirectly proves that studying PED has positive implications for CF. Although studies have shown that animal and plant-based proteins have a protective effect on CI [78], there is no specific study on the association between PED diets and CI among the population of older adults in China.

The incidence of CI in the elderly seems to be decreasing, indicating that the cohort effect of lifestyle factors is playing a role, and diet may also be a promising strategy for delaying, slowing down, or preventing CI [79–82]. A study has found that Mediterranean and DASH diets are associated with cognitive outcomes, such as lower incidence of CI and cognitive decline, or lower risk of Alzheimer’s disease [83]. Inflammation is an important mechanism of cognitive dysfunction. The Systemic Immune Inflammatory Index and Systemic Inflammatory Response Index are two blood inflammatory markers

associated with many chronic disorders, including CI. Cohort studies have shown that pro-inflammatory diets are significantly associated with CI [84]. Korean cross-sectional studies have shown that DII scores are negatively correlated with overall CF and verbal memory [85], and higher DII scores are associated with CI in women aged 65 to 79 [86]. Anti-inflammatory foods include vegetables, fruits, legumes and their products, nuts, and tea. The above dietary studies that are beneficial for CF include some anti-inflammatory foods. Consequently, it is of great significance for us to systematically construct AID and explore the relationship between AID and CI. Our study found that AID and PED patterns have a protective effect on CI; the higher the score of the two dietary patterns, the lower the probability of CI, which is very meaningful for improving and preventing CI in older adults from a dietary perspective in the future.

In addition to diet, malabsorption and some digestive problems may also be related to CI; a study has found that malabsorption of vitamin B-12 can have an impact on cognitive performance [87], and some intestinal disorders, such as celiac disorder, can also impair CF [88,89]. Furthermore, due to reduced dietary intake and poor intestinal malabsorption, the incidence of folate deficiency is high in people aged ≥ 65 years old, and population-based studies have shown that low folate levels are associated with mild CI, dementia (especially Alzheimer's disease), and depression [90]. Our research found that when the consumption of PED and AID was greater than 3 units, their protective effect on CI was more significant. That is to say, it is recommended to frequently eat more than 3 protein-rich foods and anti-inflammatory foods every day to prevent CI. According to the linear relationship proven by RCS, the greater the consumption of the two dietary patterns, the less likely they are to suffer from CI. However, considering the issues of digestion and absorption, we cannot excessively consume these diets to prevent CI. Therefore, in order to determine the more reasonable intake doses of two diets that can prevent CI while avoiding the problem of malabsorption, it is necessary to perform further exploration through more professional research on diet and digestion in future studies.

5. Limitations

To the best of our knowledge, this is the first study to systematically construct a protein-enriched diet index and anti-inflammatory diet index and explore the relationship between them and CI. However, some limitations requiring improvement still exist. Firstly, the data on CI are self-reported and may have some bias. Secondly, although we further confirmed the reliability and robustness of the research results using subgroup analysis and sensitivity analysis, cross-sectional studies are not sufficient to infer causal relationships. Therefore, in subsequent research, we could confirm the possible causal relationship between them through cohort studies. Last but not least, although we found that when the consumption of PED and AID exceeds 3 units, their protective effect on CI is more significant (owing to the linear relationship between the two dietary patterns and CI), the optimal dosage for achieving the best effect remains to be determined. Therefore, the precise dosage recommendations for the two dietary patterns need to be further demonstrated in future cohort studies.

6. Conclusions

In this study, we discovered that higher intakes of AID and PED are associated with a lower risk of CI among older adults in China, there is a linear relationship between them, and the main results are also significant under different subgroups. These findings have important implications for prevention and control of CI from a dietary and nutritional perspective in the future.

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Data Availability Statement: The CLHLS data are available at <https://opendata.pku.edu.cn/dataverse/CHADS> (accessed on 3 February 2024).

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References

1. Fang, E.F.; Xie, C.; Schenkel, J.A.; Wu, C.; Long, Q.; Cui, H.; Aman, Y.; Frank, J.; Liao, J.; Zou, H.; et al. A research agenda for ageing in China in the 21st century (2nd edition): Focusing on basic and translational research, long-term care, policy and social networks. *Ageing Res. Rev.* **2020**, *64*, 101174. [CrossRef] [PubMed]
2. Yuan, L.; Zhang, X.; Guo, N.; Li, Z.; Lv, D.; Wang, H.; Jin, J.; Wen, X.; Zhao, S.; Xu, T.; et al. Prevalence of cognitive impairment in Chinese older inpatients and its relationship with 1-year adverse health outcomes: A multi-center cohort study. *BMC Geriatr.* **2021**, *21*, 595. [CrossRef] [PubMed]
3. Tochel, C.; Smith, M.; Baldwin, H.; Gustavsson, A.; Ly, A.; Bexelius, C.; Nelson, M.; Bintener, C.; Fantoni, E.; Garre-Olmo, J.; et al. What outcomes are important to patients with mild cognitive impairment or Alzheimer's disease, their caregivers, and health-care professionals? A systematic review. *Alzheimer's Dement.* **2019**, *11*, 231–247. [CrossRef]
4. Lipnicki, D.M.; Crawford, J.D.; Dutta, R.; Thalamuthu, A.; Kochan, N.A.; Andrews, G.; Lima-Costa, M.F.; Castro-Costa, E.; Brayne, C.; Matthews, F.E.; et al. Age-related cognitive decline and associations with sex, education and apolipoprotein E genotype across ethnocultural groups and geographic regions: A collaborative cohort study. *PLoS Med.* **2017**, *14*, e1002261. [CrossRef] [PubMed]
5. Prince, M.; Bryce, R.; Albanese, E.; Wimo, A.; Ribeiro, W.; Ferri, C.P. The global prevalence of dementia: A systematic review and metaanalysis. *Alzheimer's Dement. J. Alzheimer's Assoc.* **2013**, *9*, 63–75.e2. [CrossRef] [PubMed]
6. Livingston, G.; Sommerlad, A.; Orgeta, V.; Costafreda, S.G.; Huntley, J.; Ames, D.; Ballard, C.; Banerjee, S.; Burns, A.; Cohen-Mansfield, J.; et al. Dementia prevention, intervention, and care. *Lancet* **2017**, *390*, 2673–2734. [CrossRef]
7. Ma, W.; Wu, B.; Gao, X.; Zhong, R. Association between frailty and cognitive function in older Chinese people: A moderated mediation of social relationships and depressive symptoms. *J. Affect. Disord.* **2022**, *316*, 223–232. [CrossRef]
8. Frankish, H.; Horton, R. Prevention and management of dementia: A priority for public health. *Lancet* **2017**, *390*, 2614–2615. [CrossRef] [PubMed]
9. Hu, M.; Shu, X.; Yu, G.; Wu, X.; Välimäki, M.; Feng, H. A Risk Prediction Model Based on Machine Learning for Cognitive Impairment Among Chinese Community-Dwelling Elderly People With Normal Cognition: Development and Validation Study. *J. Med. Internet Res.* **2021**, *23*, e20298. [CrossRef]
10. Jia, L.; Du, Y.; Chu, L.; Zhang, Z.; Li, F.; Lyu, D.; Li, Y.; Li, Y.; Zhu, M.; Jiao, H.; et al. Prevalence, risk factors, and management of dementia and mild cognitive impairment in adults aged 60 years or older in China: A cross-sectional study. *Lancet Public Health* **2020**, *5*, e661–e671. [CrossRef]
11. Rowe, J.W.; Kahn, R.L. Successful aging. *Gerontologist* **1997**, *37*, 433–440. [CrossRef] [PubMed]
12. Livingston, G.; Huntley, J.; Sommerlad, A.; Ames, D.; Ballard, C.; Banerjee, S.; Brayne, C.; Burns, A.; Cohen-Mansfield, J.; Cooper, C.; et al. Dementia prevention, intervention, and care: 2020 report of the Lancet Commission. *Lancet* **2020**, *396*, 413–446. [CrossRef]
13. Gallardo-Gómez, D.; Del Pozo-Cruz, J.; Noetel, M.; Álvarez-Barbosa, F.; Alfonso-Rosa, R.M.; Del Pozo Cruz, B. Optimal dose and type of exercise to improve cognitive function in older adults: A systematic review and bayesian model-based network meta-analysis of RCTs. *Ageing Res. Rev.* **2022**, *76*, 101591. [CrossRef] [PubMed]
14. Sewell, K.R.; Erickson, K.I.; Rainey-Smith, S.R.; Peiffer, J.J.; Sohrabi, H.R.; Brown, B.M. Relationships between physical activity, sleep and cognitive function: A narrative review. *Neurosci. Biobehav. Rev.* **2021**, *130*, 369–378. [CrossRef]
15. Bossers, W.J.; van der Woude, L.H.; Boersma, F.; Hortobágyi, T.; Scherder, E.J.; van Heuvelen, M.J. A 9-Week Aerobic and Strength Training Program Improves Cognitive and Motor Function in Patients with Dementia: A Randomized, Controlled Trial. *Am. J. Geriatr. Psychiatry Off. J. Am. Assoc. Geriatr. Psychiatry* **2015**, *23*, 1106–1116. [CrossRef] [PubMed]
16. Abrego-Guandique, D.M.; Bonet, M.L.; Caroleo, M.C.; Cannataro, R.; Tucci, P.; Ribot, J.; Cione, E. The Effect of Beta-Carotene on Cognitive Function: A Systematic Review. *Brain Sci.* **2023**, *13*, 1468. [CrossRef] [PubMed]
17. Caruso, G.; Torrisi, S.A.; Mogavero, M.P.; Currenti, W.; Castellano, S.; Godos, J.; Ferri, R.; Galvano, F.; Leggio, G.M.; Grosso, G.; et al. Polyphenols and neuroprotection: Therapeutic implications for cognitive decline. *Pharmacol. Ther.* **2022**, *232*, 108013. [CrossRef] [PubMed]
18. Rajaram, S.; Jones, J.; Lee, G.J. Plant-Based Dietary Patterns, Plant Foods, and Age-Related Cognitive Decline. *Adv. Nutr.* **2019**, *10*, S422–S436. [CrossRef] [PubMed]

19. Tseng, P.T.; Zeng, B.S.; Suen, M.W.; Wu, Y.C.; Correll, C.U.; Zeng, B.Y.; Kuo, J.S.; Chen, Y.W.; Chen, T.Y.; Tu, Y.K.; et al. Efficacy and acceptability of anti-inflammatory eicosapentaenoic acid for cognitive function in Alzheimer's dementia: A network meta-analysis of randomized, placebo-controlled trials with omega-3 fatty acids and FDA-approved pharmacotherapy. *Brain Behav. Immun.* **2023**, *111*, 352–364. [\[CrossRef\]](#)
20. Stavrinou, P.S.; Andreou, E.; Aphamis, G.; Pantzaris, M.; Ioannou, M.; Patrikios, I.S.; Giannaki, C.D. The Effects of a 6-Month High Dose Omega-3 and Omega-6 Polyunsaturated Fatty Acids and Antioxidant Vitamins Supplementation on Cognitive Function and Functional Capacity in Older Adults with Mild Cognitive Impairment. *Nutrients* **2020**, *12*, 325. [\[CrossRef\]](#)
21. von Schacky, C. Importance of EPA and DHA Blood Levels in Brain Structure and Function. *Nutrients* **2021**, *13*, 1074. [\[CrossRef\]](#) [\[PubMed\]](#)
22. Gou, R.; Qin, J.; Pang, W.; Cai, J.; Luo, T.; He, K.; Xiao, S.; Tang, X.; Zhang, Z.; Li, Y. Correlation between dietary patterns and cognitive function in older Chinese adults: A representative cross-sectional study. *Front. Nutr.* **2023**, *10*, 1093456. [\[CrossRef\]](#) [\[PubMed\]](#)
23. Calder, P.C.; Bosco, N.; Bourdet-Sicard, R.; Capuron, L.; Delzenne, N.; Doré, J.; Franceschi, C.; Lehtinen, M.J.; Recker, T.; Salvioli, S.; et al. Health relevance of the modification of low grade inflammation in ageing (inflammageing) and the role of nutrition. *Ageing Res. Rev.* **2017**, *40*, 95–119. [\[CrossRef\]](#) [\[PubMed\]](#)
24. Zheng, J.; Zhou, R.; Li, F.; Chen, L.; Wu, K.; Huang, J.; Liu, H.; Huang, Z.; Xu, L.; Yuan, Z.; et al. Association between dietary diversity and cognitive impairment among the oldest-old: Findings from a nationwide cohort study. *Clin. Nutr.* **2021**, *40*, 1452–1462. [\[CrossRef\]](#) [\[PubMed\]](#)
25. Zhao, N.; Smargiassi, A.; Hudson, M.; Fritzler, M.J.; Bernatsky, S. Investigating associations between anti-nuclear antibody positivity and combined long-term exposures to NO₂, O₃, and PM_{2.5} using a Bayesian kernel machine regression approach. *Environ. Int.* **2020**, *136*, 105472. [\[CrossRef\]](#) [\[PubMed\]](#)
26. Franceschi, C.; Campisi, J. Chronic inflammation (inflammaging) and its potential contribution to age-associated diseases. *J. Gerontol. Ser. A Biol. Sci. Med. Sci.* **2014**, *69* (Suppl. 1), S4–S9. [\[CrossRef\]](#) [\[PubMed\]](#)
27. Franceschi, C.; Capri, M.; Monti, D.; Giunta, S.; Olivieri, F.; Sevini, F.; Panourgia, M.P.; Invidia, L.; Celani, L.; Scurti, M.; et al. Inflammaging and anti-inflammaging: A systemic perspective on aging and longevity emerged from studies in humans. *Mech. Ageing Dev.* **2007**, *128*, 92–105. [\[CrossRef\]](#) [\[PubMed\]](#)
28. Walker, K.A.; Ficek, B.N.; Westbrook, R. Understanding the Role of Systemic Inflammation in Alzheimer's Disease. *ACS Chem. Neurosci.* **2019**, *10*, 3340–3342. [\[CrossRef\]](#)
29. Charisis, S.; Ntanasi, E.; Yannakoulia, M.; Anastasiou, C.A.; Kosmidis, M.H.; Dardiotis, E.; Gargalionis, A.N.; Patas, K.; Chatzipanagiotou, S.; Mourtzinou, I.; et al. Diet Inflammatory Index and Dementia Incidence: A Population-Based Study. *Neurology* **2021**, *97*, e2381–e2391. [\[CrossRef\]](#)
30. Tolkien, K.; Bradburn, S.; Murgatroyd, C. An anti-inflammatory diet as a potential intervention for depressive disorders: A systematic review and meta-analysis. *Clin. Nutr.* **2019**, *38*, 2045–2052. [\[CrossRef\]](#)
31. Lv, X.; Sun, S.; Wang, J.; Chen, H.; Li, S.; Hu, Y.; Yu, M.; Zeng, Y.; Gao, X.; Xu, Y.; et al. Anti-Inflammatory Dietary Diversity and Depressive Symptoms among Older Adults: A Nationwide Cross-Sectional Analysis. *Nutrients* **2022**, *14*, 5062. [\[CrossRef\]](#) [\[PubMed\]](#)
32. Valls-Pedret, C.; Sala-Vila, A.; Serra-Mir, M.; Corella, D.; de la Torre, R.; Martínez-González, M.; Martínez-Lapiscina, E.H.; Fitó, M.; Pérez-Heras, A.; Salas-Salvador, J.; et al. Mediterranean Diet and Age-Related Cognitive Decline: A Randomized Clinical Trial. *JAMA Intern. Med.* **2015**, *175*, 1094–1103. [\[CrossRef\]](#) [\[PubMed\]](#)
33. Petersson, S.D.; Philippou, E. Mediterranean Diet, Cognitive Function, and Dementia: A Systematic Review of the Evidence. *Adv. Nutr.* **2016**, *7*, 889–904. [\[CrossRef\]](#) [\[PubMed\]](#)
34. Siervo, M.; Shannon, O.M.; Llewellyn, D.J.; Stephan, B.C.; Fontana, L. Mediterranean diet and cognitive function: From methodology to mechanisms of action. *Free Radic. Biol. Med.* **2021**, *176*, 105–117. [\[CrossRef\]](#) [\[PubMed\]](#)
35. Coelho-Júnior, H.J.; Trichopoulou, A.; Panza, F. Cross-sectional and longitudinal associations between adherence to Mediterranean diet with physical performance and cognitive function in older adults: A systematic review and meta-analysis. *Ageing Res. Rev.* **2021**, *70*, 101395. [\[CrossRef\]](#)
36. Barnes, L.L.; Dhana, K.; Liu, X.; Carey, V.J.; Ventrelle, J.; Johnson, K.; Hollings, C.S.; Bishop, L.; Laranjo, N.; Stubbs, B.J.; et al. Trial of the MIND Diet for Prevention of Cognitive Decline in Older Persons. *N. Engl. J. Med.* **2023**, *389*, 602–611. [\[CrossRef\]](#) [\[PubMed\]](#)
37. Morris, M.C.; Tangney, C.C.; Wang, Y.; Sacks, F.M.; Barnes, L.L.; Bennett, D.A.; Aggarwal, N.T. MIND diet slows cognitive decline with aging. *Alzheimer's Dement. J. Alzheimer's Assoc.* **2015**, *11*, 1015–1022. [\[CrossRef\]](#) [\[PubMed\]](#)
38. Kheirouri, S.; Alizadeh, M. MIND diet and cognitive performance in older adults: A systematic review. *Crit. Rev. Food Sci. Nutr.* **2022**, *62*, 8059–8077. [\[CrossRef\]](#)
39. Berendsen, A.A.M.; Kang, J.H.; van de Rest, O.; Feskens, E.J.M.; de Groot, L.; Grodstein, F. The Dietary Approaches to Stop Hypertension Diet, Cognitive Function, and Cognitive Decline in American Older Women. *J. Am. Med. Dir. Assoc.* **2017**, *18*, 427–432. [\[CrossRef\]](#)
40. Ma, Z.; Yang, H.; Meng, G.; Zhang, Q.; Liu, L.; Wu, H.; Gu, Y.; Zhang, S.; Wang, X.; Zhang, J.; et al. Anti-inflammatory dietary pattern is associated with handgrip strength decline: A prospective cohort study. *Eur. J. Nutr.* **2023**, *62*, 3207–3216. [\[CrossRef\]](#)

41. Duchowny, K.A.; Ackley, S.F.; Brenowitz, W.D.; Wang, J.; Zimmerman, S.C.; Caunca, M.R.; Glymour, M.M. Associations Between Handgrip Strength and Dementia Risk, Cognition, and Neuroimaging Outcomes in the UK Biobank Cohort Study. *JAMA Netw. Open* **2022**, *5*, e2218314. [[CrossRef](#)] [[PubMed](#)]
42. Kuo, K.; Zhang, Y.R.; Chen, S.D.; He, X.Y.; Huang, S.Y.; Wu, B.S.; Deng, Y.T.; Yang, L.; Ou, Y.N.; Guo, Y.; et al. Associations of grip strength, walking pace, and the risk of incident dementia: A prospective cohort study of 340212 participants. *Alzheimer's Dement. J. Alzheimer's Assoc.* **2023**, *19*, 1415–1427. [[CrossRef](#)] [[PubMed](#)]
43. Li, R.; Zhan, W.; Huang, X.; Zhang, L.; Zhang, Z.; Zhou, M.; Wang, Z.; Ma, Y. The Relationship Between Mild Cognitive Impairment and Anti-Inflammatory/Pro-Inflammatory Nutrients in the Elderly in Northern China: A Bayesian Kernel Machine Regression Approach. *J. Inflamm. Res.* **2022**, *15*, 325–339. [[CrossRef](#)] [[PubMed](#)]
44. Wu, G. Dietary protein intake and human health. *Food Funct.* **2016**, *7*, 1251–1265. [[CrossRef](#)] [[PubMed](#)]
45. La Rue, A.; Koehler, K.M.; Wayne, S.J.; Chiulli, S.J.; Haaland, K.Y.; Garry, P.J. Nutritional status and cognitive functioning in a normally aging sample: A 6-y reassessment. *Am. J. Clin. Nutr.* **1997**, *65*, 20–29. [[CrossRef](#)] [[PubMed](#)]
46. van der Zwaluw, N.L.; van de Rest, O.; Tieland, M.; Adam, J.J.; Hiddink, G.J.; van Loon, L.J.; de Groot, L.C. The impact of protein supplementation on cognitive performance in frail elderly. *Eur. J. Nutr.* **2014**, *53*, 803–812. [[CrossRef](#)]
47. Roschel, H.; Gualano, B.; Ostojic, S.M.; Rawson, E.S. Creatine Supplementation and Brain Health. *Nutrients* **2021**, *13*, 586. [[CrossRef](#)]
48. Elechi, J.O.G.; Guandique, D.M.A.; Cannataro, R. Creatine in Cognitive Performance: A Commentary. *Curr. Mol. Pharmacol.* **2024**, *17*, e18761429272915. [[CrossRef](#)]
49. Sakurai, K.; Okada, E.; Anzai, S.; Tamura, R.; Shiraishi, I.; Inamura, N.; Kobayashi, S.; Sato, M.; Matsumoto, T.; Kudo, K.; et al. Protein-Balanced Dietary Habits Benefit Cognitive Function in Japanese Older Adults. *Nutrients* **2023**, *15*, 770. [[CrossRef](#)]
50. Li, Y.; Li, S.; Wang, W.; Zhang, D. Association between Dietary Protein Intake and Cognitive Function in Adults Aged 60 Years and Older. *J. Nutr. Health Aging* **2020**, *24*, 223–229. [[CrossRef](#)]
51. O'Neill, R.F.; Brennan, L.; Prinelli, F.; Sergi, G.; Trevisan, C.; De Groot, L.; Volkert, D.; Maggi, S.; Noale, M.; Conti, S.; et al. PROtein enriched MEDiterranean diet to combat undernutrition and promote healthy neuroCOgnitive ageing in older adults: The PROMED-COG consortium project. *Nutr. Bull.* **2022**, *47*, 356–365. [[CrossRef](#)] [[PubMed](#)]
52. Zhang, Y.; Chen, J.; Qiu, J.; Li, Y.; Wang, J.; Jiao, J. Intakes of fish and polyunsaturated fatty acids and mild-to-severe cognitive impairment risks: A dose-response meta-analysis of 21 cohort studies. *Am. J. Clin. Nutr.* **2016**, *103*, 330–340. [[CrossRef](#)] [[PubMed](#)]
53. Talaei, M.; Feng, L.; Yuan, J.M.; Pan, A.; Koh, W.P. Dairy, soy, and calcium consumption and risk of cognitive impairment: The Singapore Chinese Health Study. *Eur. J. Nutr.* **2020**, *59*, 1541–1552. [[CrossRef](#)] [[PubMed](#)]
54. Ortega, R.M.; Requejo, A.M.; Andrés, P.; López-Sobaler, A.M.; Quintas, M.E.; Redondo, M.R.; Navia, B.; Rivas, T. Dietary intake and cognitive function in a group of elderly people. *Am. J. Clin. Nutr.* **1997**, *66*, 803–809. [[CrossRef](#)] [[PubMed](#)]
55. Velho, S.; Marques-Vidal, P.; Baptista, F.; Camilo, M.E. Dietary intake adequacy and cognitive function in free-living active elderly: A cross-sectional and short-term prospective study. *Clin. Nutr.* **2008**, *27*, 77–86. [[CrossRef](#)] [[PubMed](#)]
56. An, R.; Liu, G.G. Cognitive impairment and mortality among the oldest-old Chinese. *Int. J. Geriatr. Psychiatry* **2016**, *31*, 1345–1353. [[CrossRef](#)] [[PubMed](#)]
57. Zhang, Z. Gender differentials in cognitive impairment and decline of the oldest old in China. *J. Gerontol. Ser. B Psychol. Sci. Soc. Sci.* **2006**, *61*, S107–S115. [[CrossRef](#)] [[PubMed](#)]
58. Plassman, B.L.; Williams, J.W., Jr.; Burke, J.R.; Holsinger, T.; Benjamin, S. Systematic review: Factors associated with risk for and possible prevention of cognitive decline in later life. *Ann. Intern. Med.* **2010**, *153*, 182–193. [[CrossRef](#)] [[PubMed](#)]
59. Miyawaki, C.E.; Liu, M. Gender differences in cognitive impairment among the old and the oldest-old in China. *Geriatr. Gerontol. Int.* **2019**, *19*, 586–592. [[CrossRef](#)]
60. Chen, H.; Zhang, X.; Feng, Q.; Zeng, Y. The Effects of “Diet-Smoking-Gender” Three-Way Interactions on Cognitive Impairment among Chinese Older Adults. *Nutrients* **2022**, *14*, 2144. [[CrossRef](#)]
61. Baker, A.H.; Wardle, J. Sex differences in fruit and vegetable intake in older adults. *Appetite* **2003**, *40*, 269–275. [[CrossRef](#)] [[PubMed](#)]
62. D'Amico, D.; Parrott, M.D.; Greenwood, C.E.; Ferland, G.; Gaudreau, P.; Belleville, S.; Laurin, D.; Anderson, N.D.; Kergoat, M.J.; Morais, J.A.; et al. Sex differences in the relationship between dietary pattern adherence and cognitive function among older adults: Findings from the NuAge study. *Nutr. J.* **2020**, *19*, 58. [[CrossRef](#)] [[PubMed](#)]
63. Liu, J.; Shang, S.; Li, P.; Deng, M.; Chen, C.; Jiang, Y.; Dang, L.; Qu, Q. Association between current smoking and cognitive impairment depends on age: A cross-sectional study in Xi'an, China. *Med. Clin.* **2017**, *149*, 203–208. [[CrossRef](#)]
64. Chen, J.C.; Espeland, M.A.; Brunner, R.L.; Lovato, L.C.; Wallace, R.B.; Leng, X.; Phillips, L.S.; Robinson, J.G.; Kotchen, J.M.; Johnson, K.C.; et al. Sleep duration, cognitive decline, and dementia risk in older women. *Alzheimer's Dement. J. Alzheimer's Assoc.* **2016**, *12*, 21–33. [[CrossRef](#)] [[PubMed](#)]
65. Ma, Y.; Liang, L.; Zheng, F.; Shi, L.; Zhong, B.; Xie, W. Association Between Sleep Duration and Cognitive Decline. *JAMA Netw. Open* **2020**, *3*, e2013573. [[CrossRef](#)]
66. Keil, S.A.; Schindler, A.G.; Wang, M.X.; Piantino, J.; Silbert, L.C.; Elliott, J.E.; Werhane, M.L.; Thomas, R.G.; Willis, S.; Lim, M.M.; et al. Longitudinal Sleep Patterns and Cognitive Impairment in Older Adults. *JAMA Netw. Open* **2023**, *6*, e2346006. [[CrossRef](#)]

67. Muhammad, T. Life course rural/urban place of residence, depressive symptoms and cognitive impairment among older adults: Findings from the Longitudinal Aging Study in India. *BMC Psychiatry* **2023**, *23*, 391. [\[CrossRef\]](#) [\[PubMed\]](#)
68. Lee, M.Y.; Huang, X.; Hilal, S. Association between Marital Status and Cognitive Impairment in a Multi-Ethnic Asian Population. *Neuroepidemiology* **2024**, 1–9. [\[CrossRef\]](#) [\[PubMed\]](#)
69. Liu, H.; Zhang, Y.; Burgard, S.A.; Needham, B.L. Marital status and cognitive impairment in the United States: Evidence from the National Health and Aging Trends Study. *Ann. Epidemiol.* **2019**, *38*, 28–34.e2. [\[CrossRef\]](#)
70. Zhu, X.; Luchetti, M.; Aschwanden, D.; Sesker, A.A.; Stephan, Y.; Sutin, A.R.; Terracciano, A. Satisfaction With Life and Risk of Dementia: Findings From the Korean Longitudinal Study of Aging. *J. Gerontol. Ser. B Psychol. Sci. Soc. Sci.* **2022**, *77*, 1831–1840. [\[CrossRef\]](#)
71. Ding, G.; Zhao, X.; Wang, Y.; Song, D.; Chen, D.; Deng, Y.; Xing, W.; Dong, H.; Zhou, Y.; Li, D.; et al. Evaluation of the relationship between cognitive impairment and suboptimal health status in a northern Chinese population: A cross-sectional study. *J. Glob. Health* **2020**, *10*, 010804. [\[CrossRef\]](#)
72. Scarmeas, N.; Anastasiou, C.A.; Yannakoulia, M. Nutrition and prevention of cognitive impairment. *Lancet Neurol.* **2018**, *17*, 1006–1015. [\[CrossRef\]](#)
73. Cherbuin, N.; Anstey, K.J. The Mediterranean Diet is Not Related to Cognitive Change in a Large Prospective Investigation: The PATH Through Life Study. *Am. J. Geriatr. Psychiatry* **2012**, *20*, 635–639. [\[CrossRef\]](#)
74. Laitinen, M.H.; Ngandu, T.; Rovio, S.; Helkala, E.-L.; Uusitalo, U.; Viitanen, M.; Nissinen, A.; Tuomilehto, J.; Soininen, H.; Kivipelto, M. Fat Intake at Midlife and Risk of Dementia and Alzheimer’s Disease: A Population-Based Study. *Dement. Geriatr. Cogn. Disord.* **2006**, *22*, 99–107. [\[CrossRef\]](#)
75. Jayedi, A.; Shab-Bidar, S.; Eimeri, S.; Djafarian, K. Fish consumption and risk of all-cause and cardiovascular mortality: A dose–response meta-analysis of prospective observational studies. *Public Health Nutr.* **2018**, *21*, 1297–1306. [\[CrossRef\]](#)
76. Van De Rest, O.; Wang, Y.; Barnes, L.L.; Tangney, C.; Bennett, D.A.; Morris, M.C. APOE e4 and the associations of seafood and long-chain omega-3 fatty acids with cognitive decline. *Neurology* **2016**, *86*, 2063–2070. [\[CrossRef\]](#)
77. Ozawa, M.; Ohara, T.; Ninomiya, T.; Hata, J.; Yoshida, D.; Mukai, N.; Nagata, M.; Uchida, K.; Shirota, T.; Kitazono, T.; et al. Milk and Dairy Consumption and Risk of Dementia in an Elderly Japanese Population: The Hisayama Study. *J. Am. Geriatr. Soc.* **2014**, *62*, 1224–1230. [\[CrossRef\]](#) [\[PubMed\]](#)
78. Xu, X.; Yin, Y.; Niu, L.; Yang, X.; Du, X.; Tian, Q. Association between Changes in Protein Intake and Risk of Cognitive Impairment: A Prospective Cohort Study. *Nutrients* **2022**, *15*, 2. [\[CrossRef\]](#)
79. Milte, C.M.; McNaughton, S.A. Dietary patterns and successful ageing: A systematic review. *Eur. J. Nutr.* **2016**, *55*, 423–450. [\[CrossRef\]](#)
80. Matthews, F.E.; Arthur, A.; Barnes, L.E.; Bond, J.; Jagger, C.; Robinson, L.; Brayne, C. A two-decade comparison of prevalence of dementia in individuals aged 65 years and older from three geographical areas of England: Results of the Cognitive Function and Ageing Study I and II. *Lancet* **2013**, *382*, 1405–1412. [\[CrossRef\]](#)
81. Christensen, K.; Thinggaard, M.; Oksuzyan, A.; Steenstrup, T.; Andersen-Ranberg, K.; Jeune, B.; McGue, M.; Vaupel, J.W. Physical and cognitive functioning of people older than 90 years: A comparison of two Danish cohorts born 10 years apart. *Lancet* **2013**, *382*, 1507–1513. [\[CrossRef\]](#)
82. Qiu, C.; von Strauss, E.; Bäckman, L.; Winblad, B.; Fratiglioni, L. Twenty-year changes in dementia occurrence suggest decreasing incidence in central Stockholm, Sweden. *Neurology* **2013**, *80*, 1888–1894. [\[CrossRef\]](#)
83. Allès, B.; Samieri, C.; Féart, C.; Jutand, M.A.; Laurin, D.; Barberger-Gateau, P. Dietary patterns: A novel approach to examine the link between nutrition and cognitive function in older individuals. *Nutr. Res. Rev.* **2012**, *25*, 207–222. [\[CrossRef\]](#)
84. Kesse-Guyot, E.; Assmann, K.E.; Andreeva, V.A.; Touvier, M.; Neufcourt, L.; Shivappa, N.; Hébert, J.R.; Wirth, M.D.; Hercberg, S.; Galan, P.; et al. Long-term association between the dietary inflammatory index and cognitive functioning: Findings from the SU.VI.MAX study. *Eur. J. Nutr.* **2017**, *56*, 1647–1655. [\[CrossRef\]](#)
85. Shin, D.; Kwon, S.C.; Kim, M.H.; Lee, K.W.; Choi, S.Y.; Shivappa, N.; Hébert, J.R.; Chung, H.K. Inflammatory potential of diet is associated with cognitive function in an older adult Korean population. *Nutrients* **2018**, *55–56*, 56–62. [\[CrossRef\]](#)
86. Hayden, K.M.; Beavers, D.P.; Steck, S.E.; Hebert, J.R.; Tabung, F.K.; Shivappa, N.; Casanova, R.; Manson, J.E.; Padula, C.B.; Salmoirago-Blotcher, E.; et al. The association between an inflammatory diet and global cognitive function and incident dementia in older women: The Women’s Health Initiative Memory Study. *Alzheimer’s Dement. J. Alzheimer’s Assoc.* **2017**, *13*, 1187–1196. [\[CrossRef\]](#)
87. Samson, M.E.; Yeung, L.F.; Rose, C.E.; Qi, Y.P.; Taylor, C.A.; Crider, K.S. Vitamin B-12 malabsorption and renal function are critical considerations in studies of folate and vitamin B-12 interactions in cognitive performance: NHANES 2011–2014. *Am. J. Clin. Nutr.* **2022**, *116*, 74–85. [\[CrossRef\]](#)
88. Makhlof, S.; Messelmani, M.; Zaouali, J.; Mrissa, R. Cognitive impairment in celiac disease and non-celiac gluten sensitivity: Review of literature on the main cognitive impairments, the imaging and the effect of gluten free diet. *Acta Neurol. Belg.* **2018**, *118*, 21–27. [\[CrossRef\]](#)

89. Croall, I.D.; Tooth, C.; Venneri, A.; Poyser, C.; Sanders, D.S.; Hoggard, N.; Hadjivassiliou, M. Cognitive Impairment in Coeliac Disease with Respect to Disease Duration and Gluten-Free Diet Adherence: A Pilot Study. *Nutrients* **2020**, *12*, 2028. [[CrossRef](#)]
90. Araújo, J.R.; Martel, F.; Borges, N.; Araújo, J.M.; Keating, E. Folate and aging: Role in mild cognitive impairment, dementia and depression. *Ageing Res. Rev.* **2015**, *22*, 9–19. [[CrossRef](#)]

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