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**Dietary flavonoid intake and cognitive performance in older adults with
Alzheimer's type dementia**

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Dietary flavonoid intake and cognitive performance in older adults with Alzheimer's type dementia

Abstract

Background: Dietary flavonoid intake and intake of flavonoid subclasses has been associated with improved cognitive performance. However, the association between flavonoid intake and cognitive performance in older adults with Alzheimer's type dementia has not been investigated. **Objectives:** To estimate dietary total flavonoid intake and intake of flavonoid subclasses in older adults with Alzheimer's type dementia and assess the relationship of flavonoid intake with measures of cognition. **Design:** Cross sectional analysis. Setting: Community dwelling older adults in NSW, Australia. **Participants:** Older adults (+65y) with mild to moderate dementia (n=49). **Measurements:** A 24h diet recall was collected with help from a carer and used to estimate flavonoid intake. A battery of cognitive tasks assessed cognitive performance of several cognitive domains. **Results:** Pearson and spearman correlation coefficients identified an association between flavonoid intake and executive function ($r=0.319$, $p=0.025$). After controlling for depression, the relationship was reduced. **Conclusion:** The identified association between cognitive functioning, depression and flavonoid intake in older adults with Alzheimer's type dementia warrants further research in a larger sample.

Disciplines

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ORIGINAL RESEARCH

DIETARY FLAVONOID INTAKE AND COGNITIVE PERFORMANCE IN OLDER ADULTS WITH ALZHEIMER'S TYPE DEMENTIA

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Abstract: *Background:* Dietary flavonoid intake and intake of flavonoid subclasses has been associated with improved cognitive performance. However, the association between flavonoid intake and cognitive performance in older adults with Alzheimer's type dementia has not been investigated. *Objectives:* To estimate dietary total flavonoid intake and intake of flavonoid subclasses in older adults with Alzheimer's type dementia and assess the relationship of flavonoid intake with measures of cognition. *Design:* Cross sectional analysis. *Setting:* Community dwelling older adults in NSW, Australia. *Participants:* Older adults (+65y) with mild to moderate dementia (n=49). *Measurements:* A 24h diet recall was collected with help from a carer and used to estimate flavonoid intake. A battery of cognitive tasks assessed cognitive performance of several cognitive domains. *Results:* Pearson and spearman correlation coefficients identified an association between flavonoid intake and executive function (r=0.319, p=0.025). After controlling for depression, the relationship was reduced. *Conclusion:* The identified association between cognitive functioning, depression and flavonoid intake in older adults with Alzheimer's type dementia warrants further research in a larger sample.

Key words: Flavonoid, cognition, dementia.

Introduction

Flavonoids are naturally occurring plant-based phytochemicals, which are abundant in the human diet. The structure and sources of flavonoids has been well established (1, 2). Flavonoids are a subclass of polyphenols and encompass a wide group of compounds that are divided into six major classes: anthocyanidins, flavanols, flavanones, flavones, flavonols and isoflavones (1). A growing body of evidence suggests that flavonoids are non-nutritive bioactive compounds, which significantly contribute to the antioxidant activity of individual fruits and vegetables and are consequently credited with the observed health benefits (3). Flavonoids are widespread across many food sources and are found in particularly high concentrations in fruits and vegetables, wine, tea, cocoa, and soy (4). In older Australian adults, major dietary sources of flavan-3-ols and flavonols are tea (black and green) and apples; flavanones are provided by citrus fruits and juices; flavones are consumed through parsley and tomatoes; and berries and red wine provide anthocyanins (5).

Isoflavones are largely provided by the consumption of soy-based products (6).

Total dietary flavonoid consumption has been associated with improved cognitive performance and a preservation of cognitive function with ageing (7-9). More focussed research on specific flavonoid subclasses has revealed that the sub-groups flavanols, anthocyanins and flavanones may provide the most beneficial effects in the area of neuroprotection (10). There has been a particular focus in animal models, on the provision of anthocyanin rich foods, such as blueberries, when investigating cognitive outcomes, with promising results (11-13). Preliminary human trials have also begun to link anthocyanin-rich food consumption with improvements in cognitive outcomes (14-18). A high consumption of total dietary flavonoids has also been linked with a reduced risk of developing a neurodegenerative disease, such as dementia (19). However, the association between flavonoid intake and cognitive performance in older adults with Alzheimer's type dementia has not been investigated to date.

The mechanisms by which flavonoids provide neuroprotection are not well elucidated and the evidence remains largely pre-clinical. This is partially related to the quick and extensive metabolism of dietary flavonoids into metabolites with differing and largely unknown biological activities (1). In animal models,

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specific flavonoids including the flavanols catechin and epigallocatechin gallate (20) have been shown to reduce neuroinflammation and scavenge free radicals within the brain (2), a function which has been related to neuroprotection. Additionally, flavonoids, including the flavanone hesperitin have been shown to provide a range of positive neuronal effects to limit neurodegeneration, including a potential to protect neurons against injury and improve neuronal morphology (2, 21). Flavonoid supplementation has also been shown to up-regulate processes that promote learning, memory and cognition (1). Flavonol-rich cocoa consumption has shown to increase cerebrovascular blood flow, which is associated with improved cognition (22). Presently, it is unclear how many, or if all flavonoids exert such effects (1).

Evidence is mounting for the positive protective effects of flavonoid consumption for the development of dementia (7, 19, 23), including consumption of flavonoid rich fruits and vegetables, juices and wine (24, 25). However, it is also important to investigate the relationship between flavonoid consumption in older adults living with dementia and cognitive performance outcomes. This evidence could be utilized to indicate if dietary interventions with flavonoids may be warranted. It is well documented that older adults with dementia develop eating difficulties, resulting in low food intake, as dementia progresses (26). This can be related to a myriad of difficulties including dysphasia, lack of appetite, confusion about the need to eat and the loss of the ability to recognize food (27). The total dietary intake and significant sources of dietary flavonoids in older adults living with dementia has not been assessed and may differ from estimations for healthy adults aged 65+ yrs.

The aim of this study was to estimate the total dietary intake and main sources of flavonoids in older Australian adults with Alzheimer's type dementia and to assess the relationship between dietary flavonoid intake and cognitive performance.

Methods

This study was approved by the University of Wollongong and Illawarra Shoalhaven Local Health District Human Research Ethics Committee (HE11/175) and complied with current laws governing ethics in research. The current study analysed the baseline data of a randomised controlled trial, which investigated the impact of anthocyanin-rich cherry juice on cognitive outcomes in older adults with dementia (17)

Community dwelling older adults (65+ yrs) with mild to moderate Alzheimer's type dementia (as diagnosed by a hospital-based geriatrician) were recruited to the study from an outpatient clinic. Dietary data was collected by a trained nutritionist (KK) (with assistance and confirmation provided from a carer) from a single 24-h food recall. The dietary data was analysed using

the Foodworks dietary analysis package (Xyris software, version 5, 2007, Highgate Hill, QLD, Australia) (28). As dietary data relating to flavonoid content of foods is not integrated into the Xyris software, the food items were manually cross-referenced with the USDA database for the flavonoid content of selected foods (release 3.1) (29) to estimate total flavonoid consumption and the flavonoid subclasses anthocyanins, flavones, flavanones, flavonols and flavan-3-ols. A Mini Nutritional Assessment (30) was conducted as a measure of malnutrition and Lawton's Instrumental Activities of Daily Living Scale (31) measured functional ability. Anthropometric and demographic information, including education, was collected (Table 1), with assistance from a guardian or carer as appropriate.

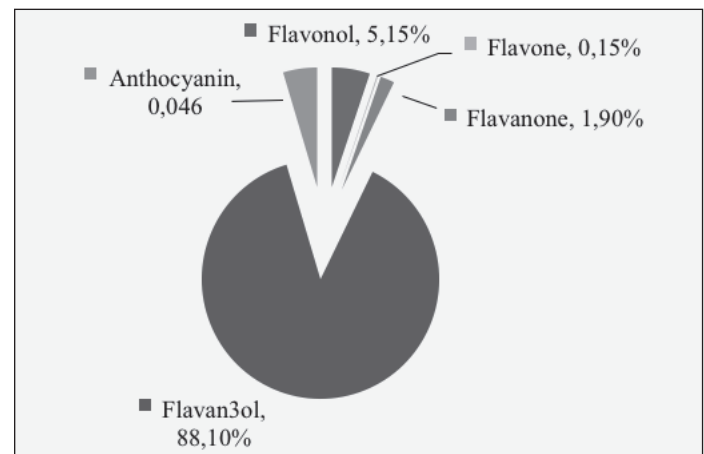
A battery of cognitive assessments was administered by a single trained researcher (KK). These included measures of mood (32), verbal learning and memory (33), working memory (34), semantic memory (35), executive function (36, 37) and short-term memory (38) (Table 2).

To assess the relationship between flavonoid intake and cognitive performance, bivariate correlations with Pearson and Spearman coefficients (as appropriate) were performed and then repeated after controlling for depression, as measured by the Geriatric Depression Scale (GDS) (32). Multiple regression analysis was performed to confirm the effect of flavonoid intake on cognitive outcomes that were identified as significant by the correlations, with age and education included in the models as covariates.

Results

Forty-nine participants volunteered to take part in the study; their anthropometric and demographic characteristics are found in Table 1.

Figure 1
Percentage contribution of flavonoid subclasses to total flavonoid intake



Analysis of the 24h dietary recall calculated total flavonoid intake to be 510.2 ± 374.8 mg/day, with a range of 5.9 – 1,524.2 mg/day. Black tea contributed

80% of total dietary intake and was the most notable dietary source of flavonoids. Other notable sources were green tea (7.5%), red wine (4.5%), with apples and oranges providing 1.7% and 1.6% respectively, when combined with their fruit juices. The dominant subclass of flavonoids was flavan-3-ols; these contributed 88% of total intake (Figure 1).

Table 1
Characteristics of study participants (n=49)

Age (years)	79.82 ± 6.9
Gender	
Male	24
Female	25
BMI (kg/m ²)	26.1 ± 3.5
Current smokers (%)	2.7%
Previous smokers (%)	40.5%
Hand grip strength (kg)	33.2 ± 6.9
Mid arm circumference (cm)	27.1 ± 4.2
Calf circumference (cm)	34.1 ± 3.6
Education (count)	
Primary	3
Secondary (year 10)	24
Secondary (year 12)	10
Tertiary	12
Mini Nutritional Assessment	24.2 ± 3.1
Lawton's Instrumental Activities of Daily Living Scale	6.2 ± 2.3

Ideal BMI for adults 65y+ is 22–27 kg/m² (13); Calf circumference and mean mid-upper arm circumference in of 29.2 in older adults; Mini Nutritional Assessment score >23.5 indicates no risk of malnutrition (14); Median maximum hand grip strength are 37.9 kg and 31.5 kg for men and women aged 50y+ (15); Lawton's IADL scale mean score 4.3 in adults with moderate dementia (11).

Total flavonoid intake was not significantly correlated with age, nutritional status or education. However, participants who displayed greater depressive symptoms (i.e. scored higher on the GDS) had a lower total flavonoid intake (r=-0.328 p=0.021).

Total flavonoid intake was significantly correlated with verbal fluency (r=0.319 p=0.025) (Table 2). Verbal fluency was also significantly correlated with the flavonoid subclasses flavonols (r=0.321 p=0.025), flavan-3-ols (r=0.323 p=0.023) and anthocyanins (r=0.298, p=0.038). No other significant associations were identified. A regression analysis was conducted to predict verbal fluency from total flavonoid intake, with age and education as covariates. Total flavonoid intake statistically significantly predicted verbal fluency score (β =0.319, p=0.025) controlling for age (β=0.042, p=0.765) and education (β=0.219, p=0.124). The overall model fit was r²=0.148.

After controlling for depression (GDS continuous score), bivariate correlations showed no significant relationship between total flavonoid intake or flavonoid subclasses was shown (Table 2).

Discussion

The mean intake of total flavonoids (510mg/day) in this sample of older adults with mild to moderate Alzheimer's type dementia is lower than the other estimations in older Australian adults, which were 683mg/day (5) (from 12 day food records) and 575mg/day (from a single 24h dietary recall) (39). However, the major dietary sources and percentage contributions from each flavonoid subclass are similar (5, 39). Black tea was the major dietary source of flavonoids (80%), and this finding highlights a limited fruit and vegetable consumption in this group.

The relationship between cognitive performance on

Table 2
Cognitive instruments, domain targeted and relationship with total flavonoid intake

Cognitive Instrument	Domain	Bivariate relationship total flavonoid intake	Bivariate relationship total flavonoid intake controlling for depression (GDS)
The Geriatric Depression scale (12)	Mood	r=-0.367, p=0.009 **	-
Rey Auditory Verbal Learning Test (16)	Verbal learning and memory	r=0.161, p=0.270	r=102, p=0.439
Self Ordered Pointing Task (17)	Working memory	r=-0.095, p=0.588	r=-0.046, p=0.760
The Boston Naming Task (18)	Semantic memory	r=0.095, p=0.514	r=-0.019, 0.899
The Verbal Fluency Task (19)	Executive function and control processes	r=0.319, p=0.025 *	r=0.191, p=0.198
Trail Making Task (20)	Executive function	r=-0.98, p=0.504	r=-0.051, p=0.773
Digit Span Backwards Task (21)	Short-term memory storage and executive control	r=0.047, p=0.747	r=-0.005, p=0.972

* Correlation is sig. at p<0.05. ** Correlation is sig. at p<0.01

the verbal fluency task (a measure of executive function), and flavonoid intake is supported by literature which suggests that executive function can be positively influenced by flavonoid supplementation and habitual flavonoid intake (40). The relationship between verbal fluency and intake of the flavonoid subclasses flavonols, flavan-3-ols and anthocyanins (but not flavanones, or flavones) may be related to the consumption of black and green tea (the major sources of flavonols and flavan-3-ols) and berries (a major source of anthocyanins), in this group. The consumption of both green and black tea has been associated with improved cognition in older adults (41), although a recent systematic review considered the role of tea in mild-cognitive impairment and dementia progression and reported that the findings were too limited to draw conclusions (42). Similarly, the consumption of anthocyanin rich strawberries and blueberries has been associated slower cognitive decline (43).

The lack of correlation between total flavonoid intake and education, nutritional status and age indicates that in this cohort these factors do not impact on total flavonoid consumption. This may be explained by tea consumption, the largest contributor to total flavonoid intake being a universally consumed item by all socioeconomic groups, and also reflects the lack of energy or micronutrients provided by this beverage, thus not impacting on nutritional status.

The positive association between depressive symptomatology (as measured by GDS) and flavonoid intake ($r=-0.367$, $p=0.009$) is interesting. The reduction in association between verbal fluency and flavonoid intake when depression was included as a covariate suggests that this relationship is confounded by the effect of depression on executive functioning. We speculate that epidemiological studies that have reported associations between flavonoid intake and cognitive outcomes, without controlling for depression, may have overestimated the strength of this relationship (7).

There is little consensus regarding which cognitive domain is impacted by flavonoid intake (40, 44) or which tool should be employed to measure this effect (44). We therefore used a battery of cognitive tests in order to investigate a wide range of domains of cognitive function, ensuring that each of the validated tests was sensitive and specific. Generalizability of the study findings is limited by the small sample size and the non representative nature of the convenience study sample. Other limitations include the inadequacy of using a single 24h dietary recall to estimate habitual dietary flavonoid intake. Aside from the limitations associated with this dietary assessment method, there are several well documented problems associated with utilising food composition databases to assign flavonoid content to selected foods to reflect dietary intake. These limitations include an incomplete list of flavonoid containing foods, non-regional specific data

(there is no Australian database for the flavonoid content of foods comprising the Australian food supply), and the inability of a dietary assessment to account for the high intra-individual variation of flavonoid absorption and metabolism (45). These limitations may result large variations in estimations of flavonoid intake (6) that differ from the true value, and may hinder our interpretation of observational data that associates flavonoid intake with specific health outcomes.

In conclusion, dietary intake of flavonoids in a sample of older adults with Alzheimer's disease was found to be somewhat lower than current Australian estimations for this age group, but the contributions of dietary sources are similar to the general older population. The identified association between cognitive functioning, depression and flavonoid intake in older adults with Alzheimer's type dementia warrants further research in a larger sample to confirm the findings and to identify whether dietary interventions may be indicated.

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Conflict of interest: None

Ethical standards: This study was approved by the University of Wollongong and Illawarra Shoalhaven Local Health District Human Research Ethics Committee (HE11/175) and complied with current laws governing ethics in research.

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