

Association between carbohydrate nutrition and prevalence of depressive symptoms in older adults

Bamini Gopinath^{1*}, Victoria M. Flood^{2,3}, George Burlutksy¹, Jimmy C. Y. Louie⁴ and Paul Mitchell¹

¹Centre for Vision Research, The Westmead Institute, The University of Sydney, Sydney, NSW 2145, Australia

²Faculty of Health Sciences, The University of Sydney, Sydney, NSW 2006, Australia

³St Vincent's Hospital, Sydney, NSW 2010, Australia

⁴School of Biological Sciences, Faculty of Science, Pok Fu Lam, University of Hong Kong, Hong Kong

(Submitted 18 July 2016 – Final revision received 13 November 2016 – Accepted 27 November 2016 – First published online 9 January 2017)

Abstract

We aimed to examine the relationship between dietary glycaemic index (GI) and glycaemic load of foods consumed, intakes of carbohydrates, sugars and fibre, and the prevalence of depressive symptoms in older adults. Data collected from 2334 participants aged 55+ years and 1952 participants aged 60+ years were analysed. Dietary information was collected using a semi-quantitative FFQ. Depressive symptoms were based on antidepressant use or either the 36-Item Short-Form Survey, which included the Mental Health Index (MHI), or the Center for Epidemiologic Studies Depression-10 Scale. Participants in the highest *v.* lowest tertile of dietary GI intake had increased odds of depressive symptoms (assessed by the MHI scale), multivariable-adjusted OR 1.55 (95% CI 1.12, 2.14). Participants in the highest compared with lowest tertile of fruit consumption had reduced odds of prevalent depressive symptoms, multivariable-adjusted OR 0.66 (95% CI 0.46, 0.95). Total fibre, vegetable fibre and breads/cereal fibre intakes were all inversely associated with the prevalence of depressive symptoms, with global *P* values of 0.03, 0.01 and 0.03, respectively. Participants in the second *v.* first tertile of vegetable consumption had 41% reduced odds of prevalent depressive symptoms, multivariable-adjusted OR 0.59 (95% CI 0.40, 0.88). We show that dietary GI and fibre intakes as well as consumption of fruits and vegetables are associated with the prevalence of depressive symptoms.

Key words: Depressive symptoms: Blue Mountains Eye Study: Carbohydrates: Glycaemic index: Fibres: Fruits: Vegetables

It has been predicted that by the year 2020 depression will be the second leading cause of burden on society among all diseases worldwide⁽¹⁾. It has been suggested that the increasing prevalence of mental disorders over recent decades might be attributed to lifestyle factors⁽²⁾. The effect of carbohydrate nutrition on health outcomes and disease has increasingly been the recent focus of research⁽³⁾. However, the link between the various aspects of carbohydrate nutrition and mental health outcomes in adults remains unclear.

Some carbohydrate-rich foods have less of an effect than others to increase blood glucose⁽⁴⁾. This property of individual foods is called the 'glycaemic index (GI)', a measure of carbohydrate quality^(4–6). Dietary glycaemic load (GL) is the product of a food's GI and total available carbohydrate content, and represents both the quantity and the quality of carbohydrates – that is, the total glycaemic impact of a portion of the food^(3,7,8). Moreover, other aspects of carbohydrate quality, such as

intakes of sugary foods and dietary fibre, are also thought to influence various health outcomes. Recent studies have tried to clarify whether some of these carbohydrate nutrition variables influence mental disorders. A recent cross-sectional study of Iranian adults⁽⁹⁾ found that there was a positive association between dietary GI intake and depression, but an inverse link between GL and mental disorders, depression and psychological distress. Similarly, cross-sectional analysis of a US cohort of homebound older adults found that dietary GI intake was positively associated with depression⁽¹⁰⁾. A cross-sectional Spanish study of institutionalised older adults⁽¹¹⁾ found an inverse link between dietary GL and prevalence of depression. Finally, one of the few longitudinal studies to assess the association between carbohydrate nutrition and depression was a US cohort study of postmenopausal women, which showed that higher GI and sugar intake were associated with an increased risk of incident depression. Conversely, higher consumption of

Abbreviations: BMES, Blue Mountains Eye Study; CES-D, Center for Epidemiologic Studies Depression; GI, glycaemic index; GL, glycaemic load; MHI, Mental Health Index.

* **Corresponding author:** Associate Professor B. Gopinath, fax +61 2 8627 3099, email bamini.gopinath@sydney.edu.au

fibre, fruits and vegetables was associated with lower risk of depression in this study⁽¹²⁾.

Given the equivocal nature of the findings on the relationship between carbohydrate nutrition and mental health in older adults, the aim of the present study was to clarify the independent association of various aspects of carbohydrate nutrition (total dietary carbohydrate, GI, GL, fibre from different food sources and total sugars) and the major carbohydrate-containing food groups (fruits, vegetables and breads/cereals) with the prevalence of depressive symptoms using a large, population-based data set.

Methods

Study population

The Blue Mountains Eye Study (BMES) is a population-based cohort study of common eye diseases and other health outcomes in a suburban Australian population located west of Sydney. Study methods and procedures have been described elsewhere⁽¹³⁾. Baseline examinations of 3654 residents aged >49 years were conducted during 1992–1994 (BMES-1, 82.4% participation rate). Surviving baseline participants were invited to attend examinations after 5 (1997–1999, BMES-2), 10 (2002–2004, BMES-3) and 15 years (2007–2009, BMES-4) at which 2334 (75.1% of survivors), 1952 (75.6% of survivors) and 1149 participants (55.4% of survivors) were re-examined, respectively, with complete data. The University of Sydney and the Western Sydney Area Human Ethics Committees approved the study, and written informed consent was obtained from all participants at each examination. All study methods were performed in accordance with the approved guidelines.

Nutritional assessment

Dietary data were collected using a 145-item, self-administered FFQ, modified for Australian diet and vernacular from an early Willett FFQ⁽¹⁴⁾, including reference portion sizes. Participants used a nine-category frequency scale to indicate the usual frequency of consuming individual food items during the past year. This FFQ included questions about the types of breakfast cereals consumed, which were used to increase the accuracy of the GI calculations. The FFQ showed reasonable agreement for carbohydrates and GI, yielding correlation coefficients of 0.55 and 0.57, respectively, and correctly classifying nearly 80% of participants within one quintile for carbohydrate intake and 74% participants within one quintile for GI, in a subset of participants who had completed 12 d of weighed food records over three seasons^(15,16).

A dietitian coded data from the FFQ into a customised database that incorporated the Australian Tables of Food Composition 1995 (NUTTAB 95) and published GI values with the scale of glucose = 100⁽¹⁷⁾. Additional GI data were obtained from the Sydney University GI Research Service online database⁽¹⁸⁾. In total, 88.9% of GI values were obtained from published values and 11.1% were interpolated from those of similar food items⁽¹⁵⁾.

An overall GI value for each participant's diet was calculated by summing the weighted GI of individual foods in the diet. The weighting was proportional to the contribution of

individual foods to total available carbohydrate intake. The GL of each food was calculated by multiplying its GI by the amount of available carbohydrate (g) per serving. Multiplication of each food's GL by frequency of consumption, summated for all food items, gave the overall dietary GL. We calculated data on total fibre intake as well as the fibre contribution from cereals, vegetables and fruits. Rolled oats and whole-grain/wholemeal bread consumption contributed predominantly to the total cereal fibre intake, that is, 46.9%. Finally, we analysed the consumption of main carbohydrate-containing food groups: vegetables, fruit, bread and cereals (comprising breakfast cereals, bread (white or other), pasta and rice).

Assessment of depressive symptoms

The Mental Health Index (MHI) component of the 36-Item Short-Form Survey (which is used to determine health-related quality of life) was administered at BMES-2 and consists of five questions regarding nervousness, happiness and low mood, feeling down, and feeling calm or peaceful. Scores were calculated as the sum of questions one to five multiplied by 25 and the result was divided by 100, that is, $\sum(\text{MH1}–\text{MH5}) \times 25/100$. Items MH3 and MH5 were coded in a reverse manner to MH1, MH2 and MH4. A cut-off score of ≤ 59 out of a total of 100 was used to define persons with significant depressive symptoms⁽¹⁹⁾ or those who used antidepressant medications.

The ten-item version of the Center for Epidemiologic Studies Depression (CES-D-10) Scale measures depressive feelings and behaviours experienced during the past week⁽²⁰⁾ and was administered at BMES-3. This modified scale has been validated in older samples and found reliable⁽²¹⁾. Measurement properties of the CES-D-10 have shown satisfactory test–re-test correlation and good predictive accuracy compared with the standard twenty-item CES-D version⁽²⁰⁾. Questions focused on mood (five items), irritability (one item), energy (two items), concentration (one item) and sleep (one item). Each of the ten items was coded on a scale of 0–3 to give a maximum of 30 points. Items 5 and 8 were coded in a reverse manner to the remaining eight items of the CES-D-10 scale. Higher scores indicate a greater burden of depressive symptoms. A cut-off score of ≥ 10 out of a total possible score of 30 was used to define participants with significant depressive symptoms⁽²⁰⁾ or use of antidepressant medications.

Assessment of covariates

A face-to-face interview with trained interviewers was conducted, and comprehensive data including information about medical histories, hearing, demographic factors, socio-economic characteristics, lifestyle and health risk behaviour such as exercise, and smoking were obtained from all participants. Participants were also asked whether they received a pension, and if so the type of pension they were receiving, for example, age, disability, veteran's or blind. Medical histories included CVD or other systemic disease and associated risk factors and medications used. A past history of angina, diabetes, myocardial infarction and stroke was determined by responses to a question: 'Has a doctor advised you that you have any of the following conditions?'. Walking difficulty or use of a cane, walker or



wheelchair was observed by a trained examiner and categorised as ‘disability in walking’. Cognitive function was assessed using the mini-mental state exam (MMSE) administered at both the baseline and the follow-up visits. MMSE scores ranged from 0 to 30⁽²²⁾, with scores <24 indicating cognitive impairment.

Statistical analysis

SAS 9.2 software (SAS Institute) was used for statistical analyses including *t* tests, χ^2 tests and logistic regression. Study factors were carbohydrate nutrition (total carbohydrates, GI, GL, sugars and fibre), and the study outcome was the prevalence of depressive symptoms as assessed by the MHI or CES-D-10 scale. Carbohydrate nutrition variables were analysed as categorical variables (tertiles). Carbohydrate nutrition variables were energy adjusted using the residual method described by Willett & Stampfer⁽²³⁾. Multivariable logistic regression models adjusted for age, sex, cognitive impairment (MMSE <24), walking disability, receiving pension, antidepressant use, previous history of stroke and arthritis were used. The logistic regression analyses are expressed as adjusted OR and 95% CI. Statistical significance was defined as *P* < 0.05.

Results

Carbohydrate nutrition and depressive symptoms as assessed by the Mental Health Index scale

At BMES-2 of the 2334 participants examined, 1918 had complete information on dietary intakes, MHI scale scores and antidepressant medication use, and hence were included in the cross-sectional analysis. Of the 1918, 275 (14.3%) had depressive symptoms. Participants with depressive symptoms compared with those without were more likely to be female, receive pension, have a walking disability and cognitive impairment, as well as have higher intakes of dietary GI and GL (online Supplementary Table S1). Table 1 shows the association between tertiles of carbohydrate nutrition variables and prevalence of depressive symptoms. After multivariable adjustment, participants in the highest tertile compared with the lowest tertile (reference group) of dietary GI intake had a 55% higher likelihood of having depressive symptoms (Table 1). Dietary intake of fibre was not associated with the prevalence of depressive symptoms (Table 2). In terms of food group consumption, the only one that showed a significant association with prevalence of depressive symptoms was fruit consumption, comparing the highest *v.* first tertile of fruit consumption, multivariable-adjusted OR 0.64 (95% CI 0.46, 0.89); global *P* = 0.04. Vegetable and breads/cereal consumption was not associated with the prevalence of depressive symptoms (data not shown).

Carbohydrate nutrition and depressive symptoms as assessed by the Center for Epidemiologic Studies Depression-10 scale

Depressive symptoms as assessed by the CES-D-10 scale were only available at BMES-3 onwards; hence, of the

Table 1. Association between carbohydrate nutrition and prevalence of depressive symptoms as assessed by the Mental Health Index scale and antidepressant use (Odds ratios and 95% confidence intervals)

Energy-adjusted variable	Prevalence of depressive symptoms		
	No. of cases/ no. at risk	OR	95% CI*
Mean dietary GI			
First tertile (≤ 54.6)	81/639	1.0 (Ref.)	
Second tertile (54.6–58.1)	79/640	0.94	0.67, 1.32
Third tertile (≥ 58.2)	115/639	1.55	1.12, 2.14
<i>P</i> _{for trend}			0.01
Global <i>P</i>			0.04
Mean dietary GL			
First tertile (≤ 126.4)	87/639	1.0 (Ref.)	
Second tertile (126.4–144.4)	84/640	0.95	0.68, 1.32
Third tertile (≥ 144.5)	84/639	1.20	0.87, 1.66
<i>P</i> _{for trend}			0.26
Global <i>P</i>			0.36
Mean total carbohydrate			
First tertile (≤ 226.7)	93/639	1.0 (Ref.)	
Second tertile (226.9–255.7)	86/640	0.85	0.61, 1.18
Third tertile (≥ 255.8)	96/639	0.98	0.71, 1.35
<i>P</i> _{for trend}			0.89
Global <i>P</i>			0.73
Mean total sugar intake			
First tertile (≤ 225.1)	94/639	1.0 (Ref.)	
Second tertile (225.3–252.9)	87/640	0.88	0.64, 1.23
Third tertile (≥ 253.0)	94/639	0.95	0.68, 1.31
<i>P</i> _{for trend}			0.77
Global <i>P</i>			0.61

GI, glycaemic index; Ref., referent values; GL, glycaemic load.
* Adjusted for age, sex, cognitive impairment (mini-mental state exam <24), walking disability, receiving pension, antidepressant use, previous history of stroke and arthritis.

1952 participants examined at BMES-3, 1504 had complete information on dietary intakes and CES-D-10 scores, and therefore were included in subsequent cross-sectional analyses. Of the 1504, 249 (16.6%) had depressive symptoms. Participants with compared with those without depressive symptoms were more likely to be female, receive pension, take antidepressants, have a walking disability, stroke, arthritis and cognitive impairment, as well as lower intake of breads/cereal fibre (online Supplementary Table S2).

Table 3 shows no significant associations between the dietary intakes of GI, GL and total carbohydrates and prevalence of depressive symptoms. However, we found that participants in the second *v.* first tertile of total fibre intake and vegetable fibre intake had a 42 and 46% reduced likelihood of having depressive symptoms, respectively (Table 4). In addition, those in the highest compared with the lowest tertile of bread/cereal fibre intake had 42% reduced odds of depressive symptoms as assessed by the CES-D-10 scale, multivariable-adjusted OR 0.58 (95% CI 0.39, 0.86). With regard to food groups, the only significant association observed was with vegetable consumption. Participants in the second tertile compared with those in the first tertile of vegetable consumption had 41% reduced odds of prevalent depressive symptoms, multivariable-adjusted OR 0.59 (95% CI 0.40, 0.88); global *P* = 0.02.

Table 2. Associations between baseline intakes of energy-adjusted dietary fibre and prevalence of depressive symptoms assessed by the Mental Health Index scale (Odds ratios and 95% confidence intervals)

Energy-adjusted variable	Prevalence of depressive symptoms		
	No. of cases/ no. at risk	OR	95% CI*
Mean total fibre intake			
First tertile (≤ 21.1)	95/639	1.0 (Ref.)	
Second tertile (21.2–27.5)	99/640	1.05	0.76, 1.44
Third tertile (≥ 27.6)	81/639	0.84	0.61, 1.18
$P_{\text{for trend}}$			0.30
Global P			0.59
Mean vegetable fibre intake			
First tertile (≤ 8.3)	96/639	1.0 (Ref.)	
Second tertile (8.4–11.5)	98/640	0.96	0.70, 1.32
Third tertile (≥ 11.6)	81/639	0.75	0.54, 1.04
$P_{\text{for trend}}$			0.08
Global P			0.25
Mean fruit fibre intake			
First tertile (≤ 4.9)	106/639	1.0 (Ref.)	
Second tertile (4.9–8.3)	87/640	0.81	0.59, 1.12
Third tertile (≥ 8.4)	82/639	0.76	0.55, 1.05
$P_{\text{for trend}}$			0.32
Global P			0.18
Mean bread/cereal fibre intake			
First tertile (≤ 5.1)	103/639	1.0 (Ref.)	
Second tertile (5.2–8.1)	84/640	0.82	0.59, 1.13
Third tertile (≥ 8.2)	88/639	0.84	0.61, 1.16
$P_{\text{for trend}}$			0.11
Global P			0.70

Ref., referent values.

* Adjusted for age, sex, cognitive impairment (mini-mental state exam < 24), walking disability, receiving pension, antidepressant use, previous history of stroke and arthritis.

Discussion

Our study findings contribute to the existing evidence base, which suggests that carbohydrate nutrition variables could be linked to the presence of depressive symptoms. Cross-sectional analysis showed that participants in the highest tertile of dietary GI intake compared with those in the lowest tertile had 55% increased likelihood of having depressive symptoms as assessed by the MHI scale and antidepressant use. Moreover, we found that total fibre, vegetable fibre and breads/cereals fibre intakes were inversely associated with the prevalence of depressive symptoms as assessed by the CES-D-10 scale. Consumption of vegetables and fruits appeared to have a modest, beneficial influence on the prevalence of depressive symptoms.

The independent association between higher dietary GI intake and greater odds of depressive symptoms observed in our cohort of older adults concurs with other epidemiological studies, which suggest that high-GI diets could be a risk factor for depression^(9,10,12). Given that this is an observational study, we were not able to determine the pathways by which dietary GI could influence mental well-being; however, we can hypothesise potential mechanisms. First, greater inflammation as a result of higher-GI intake has already been established⁽²⁴⁾, and increased inflammation has also been suggested as a mechanism for depression⁽²⁵⁾. Second, high-GI diets could also lead to insulin resistance⁽²⁶⁾, which has been associated with a pattern

Table 3. Association between carbohydrate nutrition and prevalence of depressive symptoms as assessed by the Center for Epidemiologic Studies Depression-10 scale (Odds ratios and 95% confidence intervals)

Energy-adjusted variable	Prevalence of depressive symptoms		
	No. of cases/ no. at risk	OR	95% CI*
Mean dietary GI			
First tertile (≤ 54.4)	86/509	1.0 (Ref.)	
Second tertile (54.5–58.2)	74/510	0.80	0.54, 1.19
Third tertile (≥ 58.3)	89/509	1.16	0.79, 1.71
$P_{\text{for trend}}$			0.49
Global P			0.18
Mean dietary GL			
First tertile (≤ 125.5)	91/509	1.0 (Ref.)	
Second tertile (125.6–142.4)	86/510	0.83	0.57, 1.21
Third tertile (≥ 142.4)	72/509	0.68	0.46, 1.01
$P_{\text{for trend}}$			0.06
Global P			0.17
Mean total carbohydrate			
First tertile (≤ 225.1)	86/509	1.0 (Ref.)	
Second tertile (225.3–252.9)	86/510	0.81	0.55, 1.20
Third tertile (≥ 253.0)	77/509	0.79	0.53, 1.17
$P_{\text{for trend}}$			0.24
Global P			0.43
Mean total sugar intake			
First tertile (≤ 225.1)	90/499	1.0 (Ref.)	
Second tertile (225.3–252.9)	79/503	0.93	0.63, 1.37
Third tertile (≥ 253.0)	80/502	0.91	0.61, 1.35
$P_{\text{for trend}}$			0.63
Global P			0.88

GI, glycaemic index; Ref., referent values; GL, glycaemic load.

* Adjusted for age, sex, energy intake, cognitive impairment (mini-mental state exam < 24), walking disability, receiving pension, antidepressant use, previous history of stroke and arthritis.

of volumetric and neurocognitive deficits, which are very similar to that found in individuals suffering from clinical depression^(12,27).

Conversely, dietary fibre intake was inversely associated with odds of depressive symptoms. Further, higher consumption of fruits and vegetables was also associated with a lower likelihood of depressive symptoms in older adults. Together, our findings confirm those of the recent US study on postmenopausal women, which found that higher consumption of fibre and vegetables was associated with lower odds of incident depression⁽¹²⁾. As inflammation has a potential role in depression⁽²⁵⁾, the protective effect of dietary fibre on mental health could be partly attributed to the avoidance of postprandial hyperglycaemic peaks⁽²⁸⁾, because recurrent postprandial hyperglycaemia leads to overproduction of reactive free radical molecules and greater release of inflammatory cytokines^(28–30). In addition, older adults with higher consumption of dietary fibre are also likely to be consuming higher amounts of nutrients that are important for a healthy nervous system, which therefore has a beneficial influence on mental health status⁽⁹⁾. Moreover, vegetables have low GI and this attribute could explain its consumption being inversely associated with the prevalence of depressive symptoms in older adults.

Our observation that total carbohydrate and total sugar intakes are not associated with depressive symptoms is in line with a US study on women, which also found non-significant

Table 4. Associations between intakes of energy-adjusted dietary fibre and prevalence of depressive symptoms as assessed by the Center for Epidemiologic Studies Depression-10 scale (Odds ratios and 95% confidence intervals)

Energy-adjusted variable	Prevalence of depressive symptoms		
	No. of cases/ no. at risk	OR	95% CI*
Mean total fibre intake			
First tertile (≤ 20.7)	97/497	1.0 (Ref.)	
Second tertile (20.8–27.3)	73/505	0.58	0.39, 0.87
Third tertile (≥ 27.3)	79/502	0.75	0.51, 1.10
$P_{\text{for trend}}$			0.18
Global P			0.03
Mean vegetable fibre intake			
First tertile (≤ 8.7)	96/498	1.0 (Ref.)	
Second tertile (8.7–11.8)	64/502	0.54	0.36, 0.81
Third tertile (≥ 11.9)	89/504	0.85	0.59, 1.25
$P_{\text{for trend}}$			0.50
Global P			0.01
Mean fruit fibre intake			
First tertile (≤ 5.3)	94/496	1.0 (Ref.)	
Second tertile (5.3–8.7)	76/508	0.92	0.62, 1.36
Third tertile (≥ 8.8)	79/500	0.84	0.57, 1.24
$P_{\text{for trend}}$			0.38
Global P			0.67
Mean bread/cereal fibre intake			
First tertile (≤ 4.4)	96/498	1.0 (Ref.)	
Second tertile (4.4–7.2)	84/503	0.81	0.56, 1.18
Third tertile (≥ 7.3)	69/503	0.58	0.39, 0.86
$P_{\text{for trend}}$			0.01
Global P			0.03

Ref., referent values.

* Adjusted for age, sex, energy intake, cognitive impairment (mini-mental state exam <24), walking disability, receiving pension, antidepressant use, previous history of stroke and arthritis.

links between carbohydrate and total sugar intake and risk of depression⁽¹²⁾. The authors of that study hypothesised that this could be due to dietary total sugars comprising a combination of various types of sugars and sugar from different sources⁽¹²⁾ and that the food source of sugar influences the GI, with higher fibre content slowing carbohydrate metabolisms and lowering the GI⁽¹²⁾. It is also interesting to note that we observed varying associations with the different carbohydrate nutrition variables based on whether depressive symptoms were assessed by the MHI or the CES-D-10 scale. Although there is a moderately high correlation between MHI and CES-D-10 scores⁽³¹⁾, it has been suggested that both instruments are likely to measure different aspects of mental health, that is, the MHI scale is a measure of psychological distress, whereas the CES-D-10 scale is a measure of current depressive symptoms. Hence, this could explain the differential observations with dietary GI and fibre intake. However, these findings could be due to chance and require validation by other population-based studies.

Strengths of our study are many, including use of a validated food questionnaire to collect dietary information and other questionnaires that permitted detailed assessment of potential confounding variables, as well as a relatively high participation rate. Nevertheless, there are limitations that need to be discussed. First, there is potential for misclassification because information on dietary intake was collected by self-report. Although random within-person variation could attenuate any

true association, the FFQ was designed to minimise this error by assessing average long-term consumption during the successive follow-up period⁽³²⁾. These repeated measurements take into account possible changes in diet with time and reduce random variation in reporting. Second, we used the CES-D-10 and MHI scales to assess depressive symptoms, which are screening tools only and not diagnostic tools for clinical depression, such as Diagnostic and Statistical Manual of Mental Disorders, 4th Edition (DSM-IV) criteria. Finally, we have attempted to adjust for a large number of confounding variables in the analyses; however, as with all observational studies, we cannot rule out the possibility that residual confounding (from unmeasured or unaccounted factors) could have influenced the analyses.

In conclusion, we found that carbohydrate nutrition variables such as dietary fibre showed a modest association with the prevalence of depressive symptoms. Given that depression is acknowledged as an important public health problem, it is important that the association between carbohydrate nutrition and mental health status in older adults is explored further, and randomised trials are needed in the future to examine whether changes to dietary GI and/or fibre content of an individual's diet could be an effective preventative strategy for depression in later life.

Acknowledgements

The Blue Mountains Eye Study was funded by the Australian National Health and Medical Research Council (grant nos 974159, 991407, 211069, 262120) and Westmead Institute. None of the funders had any role in the design, analysis or writing of this article.

The authors' responsibilities were as follows – B. G. and P. M.: study concept and design; P. M.: acquisition of data; G. B.: analysis of data; B. G., V. M. F., J. C. Y. L. and P. M.: interpretation of data; B. G.: drafting of the manuscript; B. G., V. M. F., J. C. Y. L., G. B. and P. M.: critical revision of the manuscript. All authors read and approved the final manuscript.

None of the authors has any conflicts of interest to declare.

Supplementary material

For supplementary material/s referred to in this article, please visit <https://doi.org/10.1017/S0007114516004311>

References

- Murray CJ & Lopez AD (1996) *The Global Burden of Disease: A Comprehensive Assessment of Mortality and Disability from Diseases, Injuries and Risk Factors in 1990 and Projected to 2020*, Global Burden of Disease and Injury Series, vol. 1. Boston, MA: Harvard School of Public Health.
- Sarris J, O'Neil A, Coulson CE, *et al.* (2014) Lifestyle medicine for depression. *BMC Psychiatry* **14**, 107.
- Gopinath B, Flood VM, Kifley A, *et al.* (2016) Association between carbohydrate nutrition and successful aging over 10 years. *J Gerontol A Biol Sci Med Sci* **71**, 1335–1340.
- Sacks FM, Carey VJ, Anderson CA, *et al.* (2014) Effects of high vs low glycemic index of dietary carbohydrate on cardiovascular disease risk factors and insulin sensitivity: the OmniCarb randomized clinical trial. *JAMA* **312**, 2531–2541.

5. Meigs JB, Nathan DM, D'Agostino RB Sr, *et al.* (2002) Fasting and postchallenge glycemia and cardiovascular disease risk: the Framingham Offspring Study. *Diabetes Care* **25**, 1845–1850.
6. Levitan EB, Mittleman MA, Hakansson N, *et al.* (2007) Dietary glycemic index, dietary glycemic load, and cardiovascular disease in middle-aged and older Swedish men. *Am J Clin Nutr* **85**, 1521–1526.
7. Jenkins DJ, Kendall CW, Augustin LS, *et al.* (2002) Glycemic index: overview of implications in health and disease. *Am J Clin Nutr* **76**, 266S–273S.
8. Schaumberg DA, Liu S, Seddon JM, *et al.* (2004) Dietary glycemic load and risk of age-related cataract. *Am J Clin Nutr* **80**, 489–495.
9. Haghghatdoost F, Azadbakht L, Keshteli AH, *et al.* (2016) Glycemic index, glycemic load, and common psychological disorders. *Am J Clin Nutr* **103**, 201–209.
10. Mwamburi DM, Liebson E, Folstein M, *et al.* (2011) Depression and glycemic intake in the homebound elderly. *J Affect Disord* **132**, 94–98.
11. Aparicio A, Robles F, Lopez-Sobaler AM, *et al.* (2013) Dietary glycaemic load and odds of depression in a group of institutionalized elderly people without antidepressant treatment. *Eur J Nutr* **52**, 1059–1066.
12. Gangwisch JE, Hale L, Garcia L, *et al.* (2015) High glycemic index diet as a risk factor for depression: analyses from the Women's Health Initiative. *Am J Clin Nutr* **102**, 454–463.
13. Attebo K, Mitchell P & Smith W (1996) Visual acuity and the causes of visual loss in Australia. The Blue Mountains Eye Study. *Ophthalmology* **103**, 357–364.
14. Willett WC, Sampson L, Browne ML, *et al.* (1988) The use of a self-administered questionnaire to assess diet four years in the past. *Am J Epidemiol* **127**, 188–199.
15. Barclay AW, Flood VM, Brand-Miller JC, *et al.* (2008) Validity of carbohydrate, glycaemic index and glycaemic load data obtained using a semi-quantitative food-frequency questionnaire. *Public Health Nutr* **11**, 573–580.
16. Smith W, Mitchell P, Reay EM, *et al.* (1998) Validity and reproducibility of a self-administered food frequency questionnaire in older people. *Aust N Z J Public Health* **22**, 456–463.
17. Foster-Powell K, Holt SH & Brand-Miller JC (2002) International table of glycemic index and glycemic load values: 2002. *Am J Clin Nutr* **76**, 5–56.
18. Sydney University GI Research Service (2010) The Glycemic Index. www.glycemicindex.com
19. Friedman B, Heisel M & Delavan R (2005) Validity of the SF-36 five-item Mental Health Index for major depression in functionally impaired, community-dwelling elderly patients. *J Am Geriatr Soc* **53**, 1978–1985.
20. Andresen EM, Malmgren JA, Carter WB, *et al.* (1994) Screening for depression in well older adults: evaluation of a short form of the CES-D (Center for Epidemiologic Studies Depression Scale). *Am J Prev Med* **10**, 77–84.
21. Irwin M, Artin KH & Oxman MN (1999) Screening for depression in the older adult: criterion validity of the 10-item Center for Epidemiological Studies Depression Scale (CES-D). *Arch Intern Med* **159**, 1701–1704.
22. Folstein MF, Folstein SE & McHugh PR (1975) 'Mini-mental state': a practical method for grading the cognitive state of patients for the clinician. *J Psychiatr Res* **12**, 189–198.
23. Willett W & Stampfer MJ (1986) Total energy intake: implications for epidemiologic analyses. *Am J Epidemiol* **124**, 17–27.
24. Buyken AE, Goletzke J, Joslowski G, *et al.* (2014) Association between carbohydrate quality and inflammatory markers: systematic review of observational and interventional studies. *Am J Clin Nutr* **99**, 813–833.
25. Han QQ & Yu J (2014) Inflammation: a mechanism of depression? *Neurosci Bull* **30**, 515–523.
26. Willett W, Manson J & Liu S (2002) Glycemic index, glycemic load, and risk of type 2 diabetes. *Am J Clin Nutr* **76**, 274S–280S.
27. McIntyre RS, Kenna HA, Nguyen HT, *et al.* (2010) Brain volume abnormalities and neurocognitive deficits in diabetes mellitus: points of pathophysiological commonality with mood disorders? *Adv Ther* **27**, 63–80.
28. Qi L & Hu FB (2007) Dietary glycemic load, whole grains, and systemic inflammation in diabetes: the epidemiological evidence. *Curr Opin Lipidol* **18**, 3–8.
29. Dickinson S, Hancock DP, Petocz P, *et al.* (2008) High-glycemic index carbohydrate increases nuclear factor-kappaB activation in mononuclear cells of young, lean healthy subjects. *Am J Clin Nutr* **87**, 1188–1193.
30. Buyken AE, Flood V, Empson M, *et al.* (2010) Carbohydrate nutrition and inflammatory disease mortality in older adults. *Am J Clin Nutr* **92**, 634–643.
31. Gopinath B, Wang JJ, Schneider J, *et al.* (2009) Depressive symptoms in older adults with hearing impairments: the Blue Mountains Study. *J Am Geriatr Soc* **57**, 1306–1308.
32. Tsai CJ, Leitzmann MF, Willett WC, *et al.* (2005) Glycemic load, glycemic index, and carbohydrate intake in relation to risk of cholecystectomy in women. *Gastroenterology* **129**, 105–112.