

Original Article

## Associations of Protein, Fat, and Carbohydrate Intakes With Insomnia Symptoms Among Middle-aged Japanese Workers

Eizaburo Tanaka<sup>1</sup>, Hiroshi Yatsuya<sup>1,2</sup>, Mayu Uemura<sup>1</sup>, Chiyoe Murata<sup>3</sup>,  
Rei Otsuka<sup>4</sup>, Hideaki Toyoshima<sup>5</sup>, Koji Tamakoshi<sup>6</sup>, Satoshi Sasaki<sup>7</sup>,  
Leo Kawaguchi<sup>1</sup>, and Atsuko Aoyama<sup>1</sup>

<sup>1</sup>Department of Public Health and Health Systems, Nagoya University School of Medicine, Nagoya, Japan

<sup>2</sup>Department of Public Health, Fujita Health University School of Medicine, Toyoake, Aichi, Japan

<sup>3</sup>Section of Social Participation and Support, Department of Social Science, Center for Gerontology and Social Science, National Center for Geriatrics and Gerontology, Obu, Aichi, Japan

<sup>4</sup>Preventive Nutrition, Department for Development of Preventive Medicine, National Center for Geriatrics and Gerontology, Obu, Aichi, Japan

<sup>5</sup>Health Care Center, Anjo Kosei Hospital, Anjo, Aichi, Japan

<sup>6</sup>Department of Nursing, Nagoya University School of Health Sciences, Nagoya, Japan

<sup>7</sup>Department of Social and Preventive Epidemiology, School of Public Health, The University of Tokyo, Tokyo, Japan

Received May 15, 2012; accepted October 16, 2012; released online February 16, 2013

### ABSTRACT

**Background:** Diet is a modifiable factor that may affect sleep, but the associations of macronutrient intakes with insomnia are inconsistent. We investigated the associations of protein, fat, and carbohydrate intakes with insomnia symptoms.

**Methods:** In this cross-sectional analysis of 4435 non-shift workers, macronutrient intakes were assessed by the brief-type self-administered diet history questionnaire, which requires the recall of usual intakes of 58 foods during the preceding month. Presence of insomnia symptoms, including difficulty initiating sleep (DIS), difficulty maintaining sleep (DMS), and poor quality of sleep (PQS) were self-reported. Logistic regression analysis was used to estimate odds ratios (ORs) and 95% CIs adjusted for demographic, psychological, and behavioral factors, as well as medical histories.

**Results:** Low protein intake (<16% vs ≥16% of total energy) was associated with DIS (OR 1.24, 95% CI 0.99–1.56) and PQS (OR 1.24, 95% CI 1.04–1.48), while high protein intake (≥19% vs <19% of total energy) was associated with DMS (OR 1.40, 95% CI 1.12–1.76). Low carbohydrate intake (<50% vs ≥50% of total energy) was associated with DMS (OR 1.19, 95% CI 0.97–1.45).

**Conclusions:** Protein and carbohydrate intakes in the daily diet were associated with insomnia symptoms. The causality of these associations remains to be explained.

**Key words:** difficulty initiating sleep (DIS); difficulty maintaining sleep (DMS); poor quality of sleep (PQS); cross-sectional study

### INTRODUCTION

About 30% of adults report 1 or more symptoms of insomnia.<sup>1</sup> Insomnia symptoms are associated with increased work absenteeism, accidents, hypertension, diabetes, and low quality of life.<sup>2–5</sup> The high prevalence and adverse impacts of insomnia make it a serious public health concern.

Demographic factors (female sex, older age, and low socioeconomic status),<sup>6,7</sup> psychological factors (stress and depression),<sup>8,9</sup> and behavioral factors (smoking, alcohol drinking, coffee consumption, and lack of regular

exercise)<sup>10–12</sup> have been associated with insomnia. Diet is a behavioral factor that may influence sleep.<sup>13</sup> A cross-sectional study reported an association between high fat intake and short sleep duration.<sup>14</sup> An interventional study reported that consumption of a supplement high in carbohydrate (total 130 g) 45 minutes before bedtime significantly increased rapid eye movement sleep as compared with a low carbohydrate supplement (total 47 g).<sup>15</sup> Another interventional study reported that tryptophan from dietary protein reduced sleep latency in those with long sleep latency and increased deep sleep as recorded by sleep electroencephalography.<sup>16</sup>

Address for correspondence. Hiroshi Yatsuya, Department of Public Health, Fujita Health University School of Medicine, 1-98 Dengakugakubo, Kutsukake-cho, Toyoake, Aichi 470-1192, Japan (e-mail: yatsuya@gmail.com).

Copyright © 2013 by the Japan Epidemiological Association

However, studies of the associations of usual intakes of protein, fat, and carbohydrate with insomnia symptoms are scarce and inconclusive.

To better understand the factors related to better sleep hygiene, we investigated the associations of protein, fat, and carbohydrate in the daily diet with 3 insomnia symptoms in a large population of middle-aged Japanese workers.

## METHODS

### Participants

In this cross-sectional study, we used data from the Aichi Workers' Cohort of cardiovascular disease. The cohort included 6651 civil servants (5179 men and 1472 women) in Aichi Prefecture—an urban and suburban area in central Japan. The cohort consisted of civil servants aged 35 to 66 years at recruitment in 2002 (participation rate: 62%). Most workers in the cohort were engaged in clerical work; police officers, firefighters, and public school teachers were not included in the cohort. The baseline survey in 2002 included a self-administered questionnaire on the absence or presence of insomnia symptoms, dietary habits, other lifestyle factors, psychological factors, and medical histories.

Shift workers ( $n = 1008$ ), who were mostly employed by public hospitals, were excluded from the present analysis because the behavior changes and circadian rhythm disruption associated with shift work may confound observed associations. We also excluded participants with missing values in variables for shift work ( $n = 127$ ), nutrient intake ( $n = 78$ ), and the covariates described below (medical history:  $n = 164$ , regular exercise:  $n = 153$ , coffee drinking habit:  $n = 134$ , alcohol intake:  $n = 91$ , and other variables:  $n = 38$ ). We also excluded those with extremely low or high energy intakes ( $<500$  or  $>4000$  kcal/day,  $n = 23$ ), which left 4835 for the analysis.

The objective of this study was explained to, and written informed consent was obtained from, all participants. The study protocol was approved by the Ethics Review Committee of Nagoya University School of Medicine, Nagoya, Japan.

### Insomnia symptoms and dietary habits

Insomnia symptoms were assessed in 3 domains by inquiring about the presence of the following symptoms: difficulty falling asleep (difficulty initiating sleep [DIS]), waking up often in the middle of the night (difficulty maintaining sleep [DMS]), and not feeling refreshed when waking up in the morning (poor quality of sleep [PQS]). The response choices were dichotomous (yes or no).

The participants' dietary habits during the preceding month were assessed with the brief-type self-administered diet history questionnaire (BDHQ), which is a simple version of the self-administered diet history questionnaire (DHQ). Both have been validated and used for nutritional studies in Japan.<sup>17–19</sup> Regarding the energy-adjusted nutrient intakes

determined by the BDHQ used in this study, a validation study that utilized 16-day weighed dietary records is regarded as the gold standard and reported the following correlation coefficients (males/females) between the 2 methods: 0.62/0.49 for protein, 0.70/0.65 for fat, and 0.78/0.67 for carbohydrate.<sup>19</sup>

Intakes of carbohydrate and fat were categorized according to values that corresponded to the national dietary goals for preventing lifestyle-related diseases, as recommended in Dietary Reference Intakes for Japanese, 2010.<sup>20</sup> These dietary reference values were established for primary prevention of hypertension, hyperlipidemia, stroke, and myocardial infarction and were based on epidemiologic studies and nutritional experiment studies.<sup>20</sup> Percentage of carbohydrate energy intake was classified into 3 categories ( $<50\%$ ,  $50\%$  to  $<70\%$ ,  $\geq 70\%$ ), as was fat intake ( $<20\%$ ,  $20$  to  $<25\%$ ,  $\geq 25\%$ ). The middle intake category of carbohydrate and fat is the recommended level.<sup>20</sup> As for protein, we used quartiles of % energy because there is no established standard.

### Covariates

Body mass index (BMI) was calculated by using measured height and weight. Mental well-being was assessed by a question asking whether participants considered their life meaningful and worth living. Responses were classified into 4 categories (yes, very much; yes; so-so; and not sure). Perceived stress was assessed by asking, "Do you have much stress in your daily life?". Responses were classified into 4 categories (very much, much, ordinary, and little). Smoking status was classified into 3 categories (current, former, and never); regular exercise into 2 categories (at least once a month for more than 60 minutes, and no exercise); alcohol intake estimated by BDHQ (g/day) into 5 categories (0, 1 to 20, 21 to 40, 41 to 60, and  $>60$  g/day); and coffee drinking habit was classified into 2 categories ( $\geq 1$  cup of coffee a day, and not every day). Medical histories of hypertension, hyperlipidemia, diabetes, angina, myocardial infarction, and stroke were self-reported. A positive history of hypertension, hyperlipidemia, and diabetes was defined as regular doctor's appointments, or prescribed medication, for the respective condition.

### Statistical analysis

Logistic regression analysis was used to estimate odds ratios (ORs) and 95% CIs for the 3 insomnia symptoms associated with intakes (% energy) of protein, fat, and carbohydrate adjusted for age, sex, BMI, total energy intake, perceived stress, mental well-being, smoking status, regular exercise, alcohol intake, coffee drinking habit, and medical histories. Age, BMI, and total energy intake were used as continuous variables in the analysis. Testing for linear trend was performed by applying the median intakes of protein, fat, and carbohydrate to each category and analyzing them as continuous variables. For additional analyses of thresholds

of insomnia symptoms, participants with a protein intake of 16% or more of total energy were integrated as a reference group in the analysis of DIS and PQS. Those with a protein intake less than 19% were integrated as a reference group in the analysis of DMS. Participants with a carbohydrate intake of 50% or more were integrated as the reference group in the analysis of DMS. Stratified analyses according to sex and alcohol intake were performed to examine confounding and effect modification. For the latter, we calculated *P*-values for the interaction of sex and alcohol intake with intakes of protein and carbohydrate. Two-sided *P*-values less than 0.05 were regarded as statistically significant. All computations were performed using SPSS version 18.0 for Windows.

## RESULTS

The mean age of participants was 48.3 years. Overall, 54.4% of the study subjects reported having at least 1 of the 3 insomnia symptoms. Overall, 6.7% reported having 2 symptoms, and 2.2% had all 3 symptoms. The prevalences of DIS, DMS, and PQS were 12.8%, 20.3%, and 32.4%, respectively (Table 1). Spearman correlation coefficients were 0.086 ( $P < 0.01$ ) for DIS and DMS, 0.022 ( $P = 0.12$ ) for DIS and PQS, and  $-0.079$  ( $P < 0.01$ ) for DMS and PQS. Subjects excluded due to missing values were older (50.7 vs 48.3 years), more likely to be female (22.0% vs 18.6%), and had a higher DIS prevalence (17.3% vs 12.8%).

In univariate logistic models, protein intake (% energy) was inversely associated with DIS and PQS but positively associated with DMS. Carbohydrate intake (% energy) was inversely associated with DMS (Table 2). In multivariate logistic models adjusted for covariates, protein intake remained inversely associated with PQS and positively associated with DMS. Carbohydrate intake was marginally inversely associated with DMS. A protein intake of less than 16% of total energy was associated with PQS and marginally associated with DIS, as compared with a protein intake of 16% or more. A protein intake of 19% or more was associated with DMS, as compared with a protein intake of less than 19%. A carbohydrate intake of less than 50% was marginally associated with DMS, as compared with a carbohydrate intake of 50% or more.

In analyses stratified by sex, a protein intake of less than 16% was marginally significantly associated with DIS in men (OR 1.30) but not in women (OR 1.11). A protein intake of less than 16% was significantly associated with PQS in men (OR 1.26) but not in women (OR 1.13). A protein intake of 19% or more was significantly associated with DMS in men (OR 1.41) but not in women (OR 1.40). A carbohydrate intake of less than 50% was marginally significantly associated with DMS in men (OR 1.21) but not in women (OR 1.02). A positive association of protein intake with DMS, and marginal inverse associations with DIS and PQS, were also observed

**Table 1. Characteristics of the study subjects (n = 4835)**

Variable	mean	SD
Age (years)	48.3	7.2
Body mass index	23.0	2.8
Energy (kcal/day)	1734	536
Carbohydrate (g/day)	242	88
Protein (g/day)	58	18
Fat (g/day)	42	14
	<i>n</i>	%
Male/female	3934/901	
Insomnia		
DIS	621	12.8
DMS	980	20.3
PQS	1567	32.4
Mental well-being		
Yes, very much	182	3.8
Yes	1405	29.0
So-so	2656	55.0
Not sure	592	12.2
Perceived stress		
Very much	507	10.5
Much	1892	39.1
Ordinary	2170	44.9
Little	266	5.5
Smoking status		
Current	1412	29.2
Former	1155	23.9
Never	2268	46.9
Alcohol consumption (g/day)		
0	1470	30.4
1–20	1739	36.0
21–40	763	15.8
41–60	491	10.2
>60	372	7.7
Coffee drinking habit		
Daily	3439	71.1
Regular exercise <sup>a</sup>	2727	56.4
Medical history		
Cardiovascular disease	35	0.7
Diabetes mellitus	370	7.7
Hyperlipidemia	158	3.3
Hypertension	809	16.7

DIS, difficulty initiating sleep; DMS, difficulty maintaining sleep; PQS, poor quality of sleep.

<sup>a</sup>≥60 min exercise at least once a month.

in nondrinking subjects (Table 3). There was no significant effect modification by protein intake and sex (*P* for interaction for DIS, 0.06; DMS, 0.36; PQS, 0.14), by protein intake and alcohol intake (*P* for interaction for DIS, 0.93; DMS, 0.33; PQS, 0.95), by carbohydrate intake and sex (*P* for interaction for DMS, 0.97), or by carbohydrate intake and alcohol intake (*P* for interaction for DMS, 0.31).

## DISCUSSION

We found that low protein intake was associated with DIS and PQS, while high protein intake was associated with DMS. Low carbohydrate intake was also associated with DMS.

**Table 2. Odds ratios and 95% CIs for DIS, DMS, and PQS according to intakes of protein, fat, and carbohydrate**

Variable	Odds ratios (CI)				<i>P</i> for trend <sup>b</sup>	
Protein intake categories (% energy)						
	<16	16–<19	19–<26	26≤	<16 vs 16– (ref)	<19 (ref) vs 19–
Mean energy intake (kcal/day)	1700	1642	1744	1849	1699/1745	
Number of participants	1209	1209	1209	1208	1209/3626	
DIS number of cases (%)	186 (15.4)	154 (12.7)	137 (11.3)	144 (11.9)	186/435 NA	
crude	1	0.80 (0.64–1.01)	0.70 (0.56–0.89)*	0.74 (0.59–0.94)*	0.25	1.33 (1.10–1.60)*
adjusted <sup>a</sup>	1	0.83 (0.65–1.06)	0.74 (0.54–1.01)	0.76 (0.47–1.24)		
DMS number of cases (%)	213 (17.6)	208 (17.2)	254 (21.0)	305 (25.2)	675/305	
crude	1	0.97 (0.79–1.20)	1.25 (1.01–1.53)*	1.58 (1.30–1.92)*	0.02*	1.47 (1.26–1.72)
adjusted <sup>a</sup>	1	0.98 (0.78–1.22)	1.37 (1.05–1.79)*	1.59 (1.06–2.38)*		
PQS number of cases (%)	450 (37.2)	392 (32.4)	372 (30.8)	353 (29.2)	450/1117	
crude	1	0.81 (0.68–0.96)*	0.75 (0.63–0.89)*	0.70 (0.59–0.83)*	0.49	1.33 (1.16–1.53)*
adjusted <sup>a</sup>	1	0.79 (0.66–0.95)*	0.84 (0.67–1.06)	0.87 (0.61–1.24)		
Fat intake categories (% energy)						
	<20	20–<25	25≤			
Mean energy intake (kcal/day)	1888	1724	1559			
Number of participants	1743	1633	1459			
DIS number of cases (%)	226 (13.0)	204 (12.5)	191 (13.1)			
crude	1.04 (0.85–1.28)	1	1.06 (0.85–1.30)	0.31		
adjusted <sup>a</sup>	1.06 (0.86–1.32)	1	0.94 (0.75–1.17)			
DMS number of cases (%)	380 (21.8)	321 (19.7)	279 (19.1)			
crude	1.14 (0.96–1.35)	1	0.97 (0.81–1.16)	0.38		
adjusted <sup>a</sup>	0.96 (0.80–1.15)	1	1.05 (0.87–1.27)			
PQS number of cases (%)	542 (31.1)	534 (32.7)	491 (33.7)			
crude	0.93 (0.80–1.07)	1	1.04 (0.90–1.21)	0.24		
adjusted <sup>a</sup>	0.98 (0.83–1.15)	1	0.88 (0.75–1.04)			
Carbohydrate intake categories (% energy)						
	<50	50–<70	70≤	<50 vs 50– (ref)		
Mean energy intake (kcal/day)	1693	1733	2042			
Number of participants	1104	3583	148			
DIS number of cases (%)	128 (11.6)	472 (13.2)	21 (14.2)	NA		
crude	0.86 (0.70–1.06)	1	1.09 (0.68–1.75)	0.27		
adjusted <sup>a</sup>	0.86 (0.67–1.11)	1	1.05 (0.64–1.71)			
DMS number of cases (%)	268 (24.3)	687 (19.2)	25 (16.9)	268/712		
crude	1.35 (1.15–1.59)*	1	0.86 (0.55–1.33)	0.07	1.36 (1.16–1.60)*	
adjusted <sup>a</sup>	1.18 (0.97–1.45)	1	0.85 (0.54–1.34)		1.19 (0.97–1.45)	
PQS number of cases (%)	335 (30.3)	1181 (33.0)	51 (34.5)	NA		
crude	0.89 (0.77–1.03)	1	1.07 (0.76–1.51)	0.35		
adjusted <sup>a</sup>	0.93 (0.78–1.12)	1	1.12 (0.77–1.62)			

DIS, difficulty in initiating sleep; DMS, difficulty in maintaining sleep; PQS, poor quality of sleep; ref, reference category; NA, not applicable. \**P* < 0.05.

<sup>a</sup>Odds ratios adjusted for age, sex, body mass index, mental well-being, perceived stress, smoking status, regular exercise, alcohol intake, coffee drinking habit, total energy intake, and histories of hypertension, hyperlipidemia, diabetes, and cardiovascular disease.

<sup>b</sup>Tests for linear trends used the median value in each group as a continuous variable in logistic regression.

Previous findings on the associations of protein and carbohydrate intakes with insomnia symptoms have been inconsistent. A cross-sectional study found an association between low protein and carbohydrate intakes and self-reported symptoms of insomnia (ie, DIS, DMS, and PQS).<sup>21</sup> However, another study found no such associations.<sup>22</sup> This inconsistency may be due to differences in the study population and differing definitions of insomnia. The former study investigated a small working population and used 4 questions to inquire about insomnia symptoms, including DIS, DMS, PQS, and early involuntary awakening. The latter study analyzed a large general population of older men and used a single question (“Have you had insomnia during the last 3 months?”). The present study included a large sample size of a general working population that was relatively homogeneous

in educational level, income, and work environment and evaluated a large number of covariates to control for potential confounding. In addition, 3 questions were used to assess insomnia symptoms, and a validated dietary questionnaire was used to determine macronutrient intakes.

Our findings of associations between protein intake and insomnia symptoms might seem inconsistent, since protein intake was inversely associated with DIS and PQS but positively associated with DMS. However, it is possible that some amino acids from dietary protein are associated with insomnia symptoms. For example, tryptophan is a dietary amino acid that serves as a precursor for brain serotonin, a sleep-inducing agent.<sup>16</sup> Low protein intake could lead to low availability of tryptophan and cause insomnia symptoms. Paradoxically, high protein intake could lower brain

**Table 3. Odds ratios and 95% CIs for DIS, DMS, and PQS according to intakes of protein, fat, and carbohydrate stratified by alcohol drinking status**

Alcohol drinking status	Intakes (% energy)	Insomnia symptoms	Odds ratios (95% CI) <sup>a</sup>
Drinkers ( <i>n</i> = 3365)	Protein	<16%	16%≤
		DIS	1.27 (0.90–1.80)
	PQS	1.20 (0.92–1.56)	1
	Protein	<19%	19%≤
		DMS	1
	Carbohydrate	<50%	50%≤
DMS		1.14 (0.91–1.43)	1
Nondrinkers ( <i>n</i> = 1470)	Protein	<16%	16%≤
		DIS	1.27 (0.93–1.73)
	PQS	1.24 (0.98–1.57)	1
	Protein	<19%	19%≤
		DMS	1
	Carbohydrate	<50%	50%≤
DMS		1.39 (0.88–2.19)	1

DIS, difficulty in initiating sleep; DMS, difficulty in maintaining sleep; PQS, poor quality of sleep. \* $P < 0.05$ .

<sup>a</sup>Odds ratios adjusted for age, sex, body mass index, mental well-being, perceived stress, smoking status, regular exercise, alcohol intake, coffee drinking habit, total energy intake, and histories of hypertension, hyperlipidemia, diabetes, and cardiovascular disease.

tryptophan levels since other competitive large neutral amino acids (LNAAs) contained in protein prevent tryptophan from crossing the blood brain barrier,<sup>23,24</sup> which might also cause insomnia symptoms. Another possibility is that high protein intake facilitates catecholamine syntheses by supplying the amino acid tyrosine,<sup>25</sup> which could increase alertness.<sup>26</sup> Therefore, both too little and too much protein could cause insomnia symptoms. We carried out an additional analysis of the association of protein intake with having at least 1 of the 3 insomnia symptoms and found that the association was U-shaped (quadratic  $P = 0.01$ , data not shown), which could support our hypothesis. Because the BDHQ did not allow estimation of tryptophan intake, further studies are needed to elucidate the associations between amino acid intakes, such as tryptophan and tyrosine, and insomnia symptoms in this and other populations.

Although presence of insomnia symptoms and fat intake were both inversely associated with sleep duration, no association between fat intake and insomnia symptoms was observed, which was consistent with the findings of previous studies.<sup>14,21,22,27</sup> In addition, although alcohol intake was associated with both fat intake and presence of insomnia symptoms, fat intake was not related to presence of insomnia symptom even after statistical adjustment for alcohol intake. In the future, prospective cohort studies must confirm whether such associations are present in different samples.

A previous study found that low carbohydrate intake was associated with shorter sleep duration.<sup>27</sup> Previous experimental studies suggest that increases in blood glucose level after carbohydrate intake are associated with early sleep onset and deep sleep,<sup>15,28</sup> possibly due to the specific effect of carbohydrate on tryptophan transport. By triggering secretion of insulin, carbohydrate lowers the levels of all serum amino acids except tryptophan. This results in higher serum tryptophan levels (relative to LNAAs), which allows tryptophan to cross the blood brain–barrier much more easily than other LNAAs.<sup>23,29</sup>

There are some limitations to our study. First, causality cannot be inferred from a cross-sectional study. It is possible that workers with symptoms of insomnia may have altered their nutritional intakes. However, the relation between diet and insomnia symptoms is not well-known, and it is thus unlikely that lay individuals would change their diet because of their awareness of such symptoms. Therefore, we speculate that diet influenced insomnia symptoms. Second, the percentage of subjects excluded due to missing values was relatively large (14%). Although the excluded subjects were more likely to have DIS, there were no statistical differences in macronutrient intakes between participants and nonparticipants with DIS. Therefore, we believe that our finding was not due to systematic selection of samples due to missing values. Third, the proportion of individuals with at

least 1 of the 3 insomnia symptoms (54.4%) was higher than the prevalence of insomnia in previous reports (30% to 45%), as assessed by the Pittsburgh Sleep Quality Index (PSQI), in the Japanese working population.<sup>30</sup> This may simply be due to the different definitions of insomnia used in the present and previous studies. Both the PSQI and the questionnaire used in the present study are self-reported. However, the PQSI also assesses sleep quantity and quality by inquiring about 18 items and generates a score to define insomnia.<sup>31</sup> The question items used in the present study only assess sleep quality by inquiring about the 3 separate insomnia symptoms. Therefore, caution is warranted in interpreting the differences in prevalence across studies. Another study of a Japanese working population, which used a definition similar to that in the present study (ie, presence of 1 of 2 insomnia symptoms [DIS, DMS]), found a similar prevalence (24.0% in men and 28.6% in women vs 29.3% in men and 29.4% in women in the present study).<sup>32</sup> Fourth, we relied solely on self-reports for the definition of insomnia. However, studies have indicated that self-reported insomnia shows at least moderate agreement with polysomnographic findings.<sup>33,34</sup> Fifth, the validity and reliability of the questionnaire items on symptoms of insomnia were not evaluated. Although we included 3 domains of insomnia (DIS, DMS, and PQS), which is consistent with the definition of insomnia in the International Classification of Diseases-10 Research Diagnostic Criteria (ICD-10 RDC), the questionnaire did not include frequency and duration of symptoms, which should be at least 3 times per week for at least 1 month according to the ICD-10 RDC. Therefore, presence of a symptom depended on participant perception, and there is thus a possibility of over- or under-reporting. However, such over- or under-reporting would be unrelated to nutrient intakes, and we believe that it would have weakened the association. In addition, participants who had at least 1 insomnia symptom were more likely to have high perceived stress (15.4% in those with insomnia symptoms vs 4.7% in normal sleepers,  $P < 0.01$ ), failure to complete a task in time (60.9% in those with insomnia symptoms vs 49.6% in normal sleepers,  $P < 0.01$ ), which is consistent with ICD-10 RDC criteria that require daytime dysfunction for a diagnosis of insomnia. Moreover, previous studies that used similar items to assess insomnia symptoms have shown similar prevalences of insomnia symptoms.<sup>32,35–37</sup> Sixth, depression was not assessed in this study. Although severely depressed workers would not have participated, individuals with mild to moderate depression could have been included, which might have confounded the association, since depression would alter both diet and sleep.<sup>38,39</sup> Future studies should include a validated assessment to screen for depression. Last, in women, the association of dietary habit with sleep was insignificant and attenuated, possibly due to the smaller sample size, higher prevalence of insomnia symptoms, and smaller interindividual differences in nutrients intakes as compared with men. However, the directionality of the results was similar in

men and women, and no significant effect modification was observed. Thus, we believe that our decision to combine men and women in the analysis was appropriate.

In conclusion, we found associations between a protein intake of less than 16% of total energy and both DIS and PQS, a protein intake of 19% or more and DMS, and a carbohydrate intake of less than 50% and DMS. Further prospective studies are needed to determine the causal relationships.

## ONLINE ONLY MATERIALS

Abstract in Japanese.

## ACKNOWLEDGMENTS

We are sincerely grateful to the participants, and to the healthcare personnel of the local government office. This work was supported by a Grant-in-Aid for Scientific Research (KAKENHI) from the Japan Society for the Promotion of Science (JSPS) and by grants from the Japan Arteriosclerosis Prevention Fund (JAPF).

Conflicts of interest: None declared.

## REFERENCES

1. Roth T. Insomnia: definition, prevalence, etiology, and consequences. *J Clin Sleep Med.* 2007;3(5 Suppl):S7–10.
2. Léger D, Scheuermaier K, Philip P, Paillard M, Guilleminault C. SF-36: evaluation of quality of life in severe and mild insomniacs compared with good sleepers. *Psychosom Med.* 2001;63:49–55.
3. Daley M, Morin CM, LeBlanc M, Grégoire JP, Savard J, Baillargeon L. Insomnia and its relationship to health-care utilization, work absenteeism, productivity and accidents. *Sleep Med.* 2009;10:427–38.
4. Vgontzas AN, Liao D, Bixler EO, Chrousos GP, Vela-Bueno A. Insomnia with objective short sleep duration is associated with a high risk for hypertension. *Sleep.* 2009;32:491–7.
5. Vgontzas AN, Liao D, Pejovic S, Calhoun S, Karataraki M, Bixler EO. Insomnia with objective short sleep duration is associated with type 2 diabetes: A population-based study. *Diabetes Care.* 2009;32:1980–5.
6. Sutton DA, Moldofsky H, Badley EM. Insomnia and health problems in Canadians. *Sleep.* 2001;24:665–70.
7. National Institutes of Health. National Institutes of Health State of the Science Conference statement on Manifestations and Management of Chronic Insomnia in Adults, June 13–15, 2005. *Sleep.* 2005;28:1049–57.
8. Murata C, Yatsuya H, Tamakoshi K, Otsuka R, Wada K, Toyoshima H. Psychological factors and insomnia among male civil servants in Japan. *Sleep Med.* 2007;8:209–14.
9. LeBlanc M, Mérette C, Savard J, Ivers H, Baillargeon L, Morin CM. Incidence and risk factors of insomnia in a population-based sample. *Sleep.* 2009;32:1027–37.
10. Janson C, Lindberg E, Gislason T, Elmasry A, Boman G. Insomnia in men—a 10-year prospective population based study.

- Sleep*. 2001;24:425–30.
11. Shilo L, Sabbah H, Hadari R, Kovatz S, Weinberg U, Dolev S, et al. The effects of coffee consumption on sleep and melatonin secretion. *Sleep Med*. 2002;3:271–3.
  12. Reid KJ, Baron KG, Lu B, Naylor E, Wolfe L, Zee PC. Aerobic exercise improves self-reported sleep and quality of life in older adults with insomnia. *Sleep Med*. 2010;11:934–40.
  13. Yogman MW, Zeisel SH. Diet and sleep patterns in newborn infants. *N Engl J Med*. 1983;309:1147–9.
  14. Shi Z, McEvoy M, Luu J, Attia J. Dietary fat and sleep duration in Chinese men and women. *Int J Obes (Lond)*. 2008;32:1835–40.
  15. Porter JM, Horne JA. Bed-time food supplements and sleep: effects of different carbohydrate levels. *Electroencephalogr Clin Neurophysiol*. 1981;51:426–33.
  16. Hartmann E, Spinweber CL. Sleep induced by L-tryptophan. Effect of dosages within the normal dietary intake. *J Nerv Ment Dis*. 1979;167:497–9.
  17. Sasaki S, Yanagibori R, Amano K. Self-administered diet history questionnaire developed for health education: a relative validation of the test-version by comparison with 3-day diet record in women. *J Epidemiol*. 1998;8:203–15.
  18. Sasaki S, Ushio F, Amano K, Morihara M, Todoriki O, Uehara Y, et al. Serum biomarker-based validation of a self-administered diet history questionnaire for Japanese subjects. *J Nutr Sci Vitaminol (Tokyo)*. 2000;46:285–96.
  19. Kobayashi S, Honda S, Murakami K, Sasaki S, Okubo H, Hirota N, et al. Both comprehensive and brief self-administered diet history questionnaires satisfactorily rank nutrient intakes in Japanese adults. *J Epidemiol*. 2012;22:151–9.
  20. Masato K, Satoshi S, editors. Dietary reference intakes for Japanese. Tokyo: daiichi shuppan publishing Co. Ltd.; 2010.
  21. Zadeh SS, Begum K. Comparison of nutrient intake by sleep status in selected adults in Mysore, India. *Nutr Res Pract*. 2011;5:230–5.
  22. Hakkarainen R, Partonen T, Haukka J, Virtamo J, Albanes D, Lönnqvist J. Food and nutrient intake in relation to mental wellbeing. *Nutr J*. 2004;3:14.
  23. Fernstrom JD, Faller DV. Neutral amino acids in the brain: changes in response to food ingestion. *J Neurochem*. 1978;30:1531–8.
  24. Hudson C, Hudson SP, Hecht T, MacKenzie J. Protein source tryptophan versus pharmaceutical grade tryptophan as an efficacious treatment for chronic insomnia. *Nutr Neurosci*. 2005;8:121–7.
  25. Wurtman RJ, Fernstrom JD. Control of brain monoamine synthesis by diet and plasma amino acids. *Am J Clin Nutr*. 1975;28:638–47.
  26. Lieberman HR, Spring BJ, Garfield GS. The behavioral effects of food constituents: strategies used in studies of amino acids, protein, carbohydrate and caffeine. *Nutr Rev*. 1986;44 Suppl: 61–70.
  27. Weiss A, Xu F, Storfer-Isser A, Thomas A, Ievers-Landis CE, Redline S. The association of sleep duration with adolescents' fat and carbohydrate consumption. *Sleep*. 2010;33:1201–9.
  28. Afaghi A, O'Connor H, Chow CM. High-glycemic-index carbohydrate meals shorten sleep onset. *Am J Clin Nutr*. 2007;85:426–30.
  29. Fernstrom JD, Wurtman RJ. Brain serotonin content: physiological dependence on plasma tryptophan levels. *Science*. 1971;173:149–52.
  30. Doi Y, Minowa M, Tango T. Impact and correlates of poor sleep quality in Japanese white-collar employees. *Sleep*. 2003;26:467–71.
  31. Doi Y, Minowa M, Uchiyama M, Okawa M, Kim K, Shibui K, et al. Psychometric assessment of subjective sleep quality using the Japanese version of the Pittsburgh Sleep Quality Index (PSQI-J) in psychiatric disordered and control subjects. *Psychiatry Res*. 2000;97:165–72.
  32. Doi Y, Minowa M. Gender differences in excessive daytime sleepiness among Japanese workers. *Soc Sci Med*. 2003;56:883–94.
  33. Hoch CC, Reynolds CF 3rd, Kupfer DJ, Berman SR, Houck PR, Stack JA. Empirical note: self-report versus recorded sleep in healthy seniors. *Psychophysiology*. 1987;24:293–9.
  34. Frankel BL, Coursey RD, Buchbinder R, Snyder F. Recorded and reported sleep in chronic primary insomnia. *Arch Gen Psychiatry*. 1976;33:615–23.
  35. Yokoyama E, Kaneita Y, Saito Y, Uchiyama M, Matsuzaki Y, Tamaki T, et al. Association between depression and insomnia subtypes: a longitudinal study on the elderly in Japan. *Sleep*. 2010;33:1693–702.
  36. Kaneita Y, Ohida T, Uchiyama M, Takemura S, Kawahara K, Yokoyama E, et al. The relationship between depression and sleep disturbances: a Japanese nationwide general population survey. *J Clin Psychiatry*. 2006;67:196–203.
  37. Ohida T, Osaki Y, Doi Y, Tanihata T, Minowa M, Suzuki K, et al. An epidemiologic study of self-reported sleep problems among Japanese adolescents. *Sleep*. 2004;27:978–85.
  38. Kawakami N, Roberts RE, Lee ES, Araki S. Changes in rates of depressive symptoms in a Japanese working population: life-table analysis from a 4-year follow-up study. *Psychol Med*. 1995;25:1181–90.
  39. Rogers PJ. A healthy body, a healthy mind: long-term impact of diet on mood and cognitive function. *Proc Nutr Soc*. 2001;60:135–43.