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Abstract

Objective: To assess dairy Ca intake and investigate its relationship with insomnia and other common co-morbidities including anxiety, depression and musculoskeletal pain (MSP) among university students.

Design: Cross-sectional study. Setting: University, Irbid, Jordan.

Participants: Male and female individuals (n 1000), aged 20·87 \pm 2·69 years. *Results:* Low dairy Ca intake (<1000 mg/d) was reported by 96·5 % of participants, and moderate to severe insomnia reported by 15·6 % of participants. Abnormal anxiety and depression scores were reported by 26·2 and 18·0 % of participants, respectively. MSP was reported by 42·9 % of participants. Participants with moderate to severe insomnia had lower dairy Ca, higher anxiety and depression scores and higher measures of MSP compared to participants with no insomnia (P-values < 0·05). Dairy Ca was weakly inversely correlated with Insomnia Severity Index (ISI) score, depression score and measures of MSP (P-values < 0·05). Regression analysis indicated that insomnia was predicted by low dairy Ca, anxiety, depression, MSP and smoking (P-values < 0·05). Both anxiety and depression were predicted by low dairy Ca (P-value < 0·01). MSP was predicted by increased ISI and anxiety scores (P-values < 0·05).

Conclusions: Low dairy Ca was highly prevalent and associated with insomnia and depression among university students. Individuals should be advised to increase dietary Ca intake to achieve the recommended daily amount. Further research is required to investigate a potential causal relationship between low Ca and both insomnia and its related co-morbidities.

Keywords Calcium Insomnia Anxiety Depression Musculoskeletal pain

Insomnia is a type of sleep disorder that is characterised by difficulty in sleep initiation or maintenance at night and is usually accompanied by daytime symptoms such as fatigue, sleepiness, irritability and loss of concentration⁽¹⁾. Transient insomnia occurs at a rate of approximately 30 % among adults, ⁽²⁾ while chronic insomnia is reported in approximately ~10–30 % of adults and more common in those who are divorced or widowed, elderly, female or suffer from psychiatric or medical diseases^(2,3). Chronic insomnia may adversely affect daily functioning and if prolonged may be associated with increased risk of deleterious health conditions including obesity, diabetes mellitus, hypertension and other CVD^(4,5).

There is evidence to suggest that increased dietary Ca intake, which is mainly obtained from dairy products⁽⁶⁾, is associated with improved quality of sleep and reduced insomnia^(7,8). Total plasma Ca was found to be directly related to sleep as a diurnal variation in its concentration was detected during normal sleep⁽⁹⁾. In addition, brain Ca is proposed to have a role in regulating the duration of sleep in mammals⁽¹⁰⁾. The effect of dairy products on sleep could also be due to their content of tryptophan,⁽¹¹⁾ as Ca is involved in producing melatonin, the sleep-inducing neurosecretory hormone, from tryptophan in the brain⁽⁸⁾.

Because Ca is necessary for proper musculoskeletal function⁽¹²⁾, low plasma Ca levels are usually associated

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with paresthesia in extremities, fatigue and muscle cramps and spasms that can be very painful⁽¹³⁾. These symptoms may lead to some psychiatric conditions including anxiety and depression, which can also be present due to hypocalcaemia itself⁽¹³⁾. Our previous work showed independent associations between non-cardiac musculoskeletal pain (MSP) and anxiety, as well as between anxiety and low dietary Ca intake, suggesting that anxiety and MSP could be associated with low plasma Ca levels⁽¹⁴⁾. MSP may also be associated with insomnia and the two conditions could co-occur with each of them increasing the risk of the other⁽¹⁵⁾. In addition, several studies have reported a high co-morbidity between insomnia, anxiety and depression⁽¹⁶⁻¹⁹⁾.

The aim of the current study was to assess dairy Ca intake and its relationship to insomnia and other common co-morbidities including anxiety, depression and MSP in Jordanian university students whose academic achievement is often negatively impacted by these conditions. In Jordan, dietary Ca is mainly obtained from dairy products that are available in the Jordanian market including cheese, yogurt, milk and labanah. These types of food are very popular and rich in Ca^(14,20). We hypothesised that low dairy Ca intake is highly prevalent in this population and is associated with insomnia, anxiety, depression and MSP. Understanding the relationship between Ca intake and these common ailments may be effective in improving insomnia and psychological symptoms in the study population.

Materials and methods

Study design and participants

A single institution questionnaire-based cross-sectional study was conducted between September 2019 and January 2020 at Jordan University of Science and Technology, Irbid, Jordan. The study sample consisted of 1000 enrolled undergraduate students recruited from all schools of the university. Participants were divided into two groups: 'medical students' who were specialising in medicine, dentistry, pharmacy, nursing or applied medical sciences and 'non-medical students' who were specialising in engineering, computer sciences, science and art, agriculture or architecture and design. The study procedure was ethically approved by the Institutional Review Board of the university. Written informed consent was obtained from all participants before they personally completed the questionnaires. A research assistant was available to clarify any inquiries from the participants.

Data collection

Self-reported data collection included: age, height, weight, marital status, college, academic level, smoking, number of family members, family income, exercise, as well as history of medical illness (including psychological illnesses) and

Ca supplementation during the previous month. BMI was calculated from weight and height in kg/m².

Assessment of daily dairy calcium intake

Participants were asked to report the daily frequency and type of dairy product consumed including cheese, yogurt, milk and labanah as previously described^(14,20). These are the most common dairy products available in the Jordanian market and they represent the major sources of dietary Ca in this population^(14,20). Frequency of dairy intake was defined as 0, 1, 2, 3 or more servings/d,^(14,20) where a single serve was defined as: 1 ounce of cream cheese (20 mg Ca), 1 ounce of cheddar cheese (162 mg Ca), one cup (240 ml) of milk or yogurt (300 mg Ca) and two tablespoons (2 oz) of labanah (100 mg Ca)^(14,20). Daily Ca intake was then calculated in mg/d. None of the participants reported intake of Ca supplements.

Assessment of insomnia and its severity

Insomnia was assessed using an Arabic version of the wellvalidated Insomnia Severity Index (ISI)(21) that was previously translated from English to Arabic using the standard forward-backward translation method⁽²²⁾. The Arabic translation was also pilot tested on thirty undergraduate university students who were not participants in the current study. The ISI is a self-administered seven question survey that measures severity of insomnia experienced over the previous 2 weeks. The first three questions assess the difficulty in falling asleep, difficulty in staying asleep and problems with waking up prematurely. The remaining four questions assess how the subject's satisfaction with their current sleep pattern, how noticeable it is to others that there may be sleep problem impairing the subject's quality life, whether the subject is worried about their current sleep pattern and how much the current sleep pattern is interfering with the subject's daily functioning. Each of these questions is answered by selecting a number from 0 to 4 that describes the severity of the problem and the ISI score (0-28) is calculated by adding the scores of all questions together. Participants with ISI scores from 0 to 7 are considered to have no clinically significant insomnia, participants with ISI scores from 8 to 14 are classified as having subthreshold insomnia and participants with ISI scores from 15 to 28 are considered to have clinical insomnia (moderate to severe).

Assessment of psychological symptoms (anxiety and depression)

Anxiety and depression symptoms were assessed using an Arabic version of the well-validated Hospital Anxiety and Depression Scale (HADS)⁽²³⁾. The standard forward-backward translation method was also used to translate the HADS from English to Arabic as previously described⁽¹⁴⁾. The Arabic translation was also pilot tested on thirty undergraduate university students who were not participants in the current study. The HADS consists of seven questions that





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assess anxiety and seven questions that assess depression. Each question is answered by selecting a number from 0 to 3 considering to severity. Thus, the total score for both anxiety and depression ranges from 0 to 21. Participants were considered normal (HADS-anxiety or depression scores from 0 to 7), borderline (HADS-anxiety or depression scores from 8 to 10) or abnormal (HADS-anxiety or depression scores from 11 to 21)⁽²³⁾.

Assessment of musculoskeletal pain

As previously described⁽²⁴⁾, participants were asked to report the sites and severity of chronic MSP. Areas of MSP were indicated using an illustration of the human body consisting of twelve sites including neck, shoulders, lower back, hands, wrists, palms, arms, upper legs, knees, lower legs, hips and feet. The number of painful body sites was summed. Additionally, participants were asked to report the overall intensity of MSP during the last month using a 0–10 numerical rating scale (0 reflects no MSP and 10 reflects the maximum intensity of MSP)⁽²⁴⁾.

Statistical analysis

Analysis was performed using the IBM SPSS Statistics software version 23. Data were presented as frequency (%), mean \pm sD or median (25th–75th percentiles) as appropriate. Differences in variables according to insomnia status were determined using one-way ANOVA with Tukey *post-boc* test for multiple comparisons, Kruskal–Wallis one-way ANOVA or χ^2 test as appropriate. Correlation analysis between ISI score and other continuous variables was performed using either Pearson's or Spearman's correlation test as appropriate. Predictors of insomnia, anxiety and depression were determined using multiple linear regression analysis. Predictors of MSP were determined using binary logistic regression analysis. All *P*-values were two-tailed and considered statistically significant at < 0.05.

Results

General characteristics of the participants

One thousand participants (36.7%) males and 63.3% females) were recruited, consisting of 52.4% medical and 47.6% non-medical university students, whose mean age was 20.87 ± 2.69 years. According to their academic level, 55.3% of participants were in their 1st or 2nd year (junior students) and 44.7% of participants were in their 3rd–6th year (senior students). Participants had a mean BMI of 23.75 ± 4.56 kg/m² and only 14.7% of participants were current smokers. Other characteristics of participants are reported in Table 1.

Daily dairy calcium intake

Most participants (96.5%) reported a daily dairy Ca intake that was well below the recommended daily intake of

1000 mg/d for adults, ⁽²⁵⁾ with an overall median (25th–75th percentiles) of 288·12 (130·95–487·72) mg/d.

Prevalence of insomnia and its severity

According to the ISI scoring system, 15.6% of participants suffered from moderate to severe clinical insomnia, 45.0% of participants had subthreshold insomnia and 39.4% of participants had no clinically significant insomnia. The mean ISI score for all participants was 9.31 ± 5.46 .

Prevalence of psychological symptoms (anxiety and depression)

According to the HADS scoring system, abnormal HADS-anxiety scores were reported in $26\cdot2\,\%$ of participants and abnormal HADS-depression scores were reported in $18\cdot0\,\%$ of participants. Additionally, $22\cdot5\,\%$ of participants recorded borderline HADS-anxiety scores and $24\cdot3\,\%$ of participants had borderline HADS-depression scores. The remaining participants had normal HADS-anxiety (51·3 $\,\%$) and depression (57·7 $\,\%$) scores. The mean HADS-anxiety and depression scores for all participants were $7\cdot8\pm4\cdot33$ and $7\cdot13\pm3\cdot7$, respectively.

Prevalence of musculoskeletal pain

MSP was reported by 42.9% of participants. The median (25th–75th percentiles) MSP intensity (out of 10) during the previous month was 0 (0–4) and the median (25th–75th percentiles) number of painful body sites (out of 12) was 0 (0–1.75).

Differences between study variables according to insomnia status

As shown in Table 1, moderate to severe insomnia was more prevalent in non-medical students compared with medical students (P-value = 0·04). Current smokers, participants with a history of medical illness and those who reported MSP during the previous month reported higher frequencies of moderate to severe insomnia compared with subthreshold insomnia and no clinically significant insomnia (P-values = 0·01, <0·01, <0·001, respectively). Participants with moderate to severe insomnia had higher HADS-anxiety and depression scores, higher MSP intensity during the previous month, more painful body sites and lower daily dairy Ca intake compared to participants with subthreshold insomnia and participants with no clinically significant insomnia (P-values < 0·001).

Correlation between Insomnia Severity Index score, Hospital Anxiety and Depression Scale-anxiety and depression scores, musculoskeletal pain and daily dairy calcium intake

As shown in Table 2, there was a relatively strong correlation between the ISI and HADS-anxiety scores (r = 0.43, P-value < 0.001), and moderate correlations between ISI



Table 1 Differences between variables according to insomnia status

	Total (<i>n</i> 1000)		No clinically significant insomnia (ISI score = 0-7) (n 394)		Subthreshold insomnia (ISI score = 8–14) (<i>n</i> 450)		Moderate to severe clinical insomnia (ISI score = 15–28) (<i>n</i> 156)		
	n	%	n	%	п	%	n	%	P*
Age									
Mean	20.87		20.95		20.83		21.01		0.68
SD	2.69		2.72		2.37		2.60		
Gender									
Male	367	36.7	140	35.5	165	36.7	62	39.7	0.65
Female	633	63.3	254	64.5	285	63.3	94	60.3	
Marital status							• •		
Single	978	97.8	381	96.7	445	98.9	152	97.4	0.08
Married	22	2.2	13	3.3	5	1.1	4	2.6	0 00
College			10	00	· ·		-	20	
Medical	524	52.4	214	54.3	243	54.0	67	42.9	0.04
Non-medical	476	47.6	180	45.7	207	46.0	89	57·1	0.04
	470	47.0	100	43.7	207	46.0	09	37.1	
Academic level	550	55.0	04.4	540	050	50.0	00	·	0.07
Junior students (1st and 2nd years)	553	55.3	214	54.3	253	56.2	86	55.1	0.87
Senior students (3rd–6th year)	447	44.7	180	45.7	197	43.8	70	44.9	
BMI (kg/m²)									
Mean	23.75		23.92		23.64		23.63		0.73
SD	4.56		4.62		4.54		4.50		
Smoking									
Yes	147	14.7	43	10.9	72	16⋅0	32	20.5	0.01
No	853	85.3	351	89-1	378	84.0	124	79.5	
Number of family members									
Mean	6.47		6.33		6.56		6.57		0.26
SD	1.74		1.74		1.68		1.89		
Average family income									
<500 JD	246	24.6	98	24.9	103	22.9	45	28.8	
501–1000 JD	402	40.2	162	41.1	175	38.9	65	41.7	0.31
>1000 JD	352	35.2	134	34.0	172	38.2	46	29.5	001
Regular exercise	002	03.2	104	04.0	172	00.2	40	20.0	
Yes	664	66-4	273	69.3	294	65.3	97	62.2	0.23
No	336	33.6	121	30.7	294 156	34·7	59	37·8	0.23
	330	33.0	121	30.7	100	34.7	59	37.6	
History of medical illness during previous month	400	40.0	0.5	0.0	40	0.0	00	40.0	0.04
Yes	106	10.6	35	8.9	42	9.3	29	18-6	<0.01
No	894	89.4	359	91.1	408	90.7	127	81∙4	
HADS-anxiety score (0-21)									
Mean	7⋅8		5.99†‡		8·22†§		11·18‡§		0.00
SD	4.33		3.91		3⋅88		4.30		
Normal (0-7)	513	51⋅3	274	69.5	206	45⋅8	33	21.2	
Borderline (8–10)	225	22.5	64	16.2	125	27.8	36	23.1	0.00
Abnormal (11–21)	262	26.2	56	14.2	119	26.4	87	55.8	
IADS-depression score (0–21)									
Mean	7.13		6.09†‡		7·27 ^{†§}		9·37 ^{‡§}		0.00
SD	3.7		3.36		3.48		4.08		- 00

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Table 1 Continued

	Total (<i>n</i> 1000)		No clinically significant insomnia (ISI score = 0-7) (n 394)		Subthreshold insomnia (ISI score = 8–14) (n 450)		Moderate to severe clinical insomnia (ISI score = 15–28) (n 156)		
	n	%	n	%	n	%	n	%	P*
Normal (0–7)	577	57.7	268	68.0	249	55.3	60	38.5	0.000
Borderline (8–10)	243	24.3	81	20.6	126	28.0	36	23.1	
Abnormal (11–21)	180	18.0	45	11.4	75	16.7	60	38.5	
Daily dairy Ca intake (mg/d)									
Median	288.12		310.08		277.28		219.64		0.04
25th–75th percentiles	130.95-487.72		135.56-529.76		132.79-476.48		105.14-466.69		
Complaint of MSP in any site of the body									
Yes	429	42.9	138	35.0	200	44.4	91	58.3	0.000
No	571	57⋅1	256	65.0	250	55.6	65	41.7	
MSP intensity during the previous month (0–10)									
Median	0		0		0		2		0.000
25th-75th percentiles	0–4		0–2		0–4		0–5		
Number of painful body sites (1–12)									
Median	0		0		0		1		0.000
25th-75th percentiles	0-1.75		0–1		0–2		0–2		

ISI, Insomnia Severity Index; HADS, Hospital Anxiety and Depression Score; MSP, musculoskeletal pain; JD, Jordanian Dinar.

Table 2 Correlation between ISI score and other variables*

	ISI score	Daily dairy Ca intake (mg/d)	HADS-anxiety score	HADS-depression score	MSP intensity during the previous month (0–10)	Number of painful body sites (1–12)
Age (years)	r = 0.01; $P = 0.74$	r=0.09; P<0.01	r = -0.11; $P < 0.001$	r = -0.04; $P = 0.20$	r = -0.04; $P = 0.17$	r = -0.08; $P = 0.02$
BMI (kg/m²)	r = -0.01; $P = 0.78$	r = 0.04; $P = 0.28$	r = -0.02; $P = 0.65$	r = -0.02; $P = 0.52$	r = 0.06; $P = 0.04$	r = 0.03; $P = 0.44$
Number of family members	r = 0.05; $P = 0.12$	r = -0.09; $P < 0.01$	r = 0.02; $P = 0.46$	r = 0.07; $P = 0.02$	r = 0.01; $P = 0.77$	r = -0.02; $P = 0.55$
ISI score	_	r = -0.10; $P < 0.01$	r = 0.43; $P < 0.001$	r = 0.27; $P < 0.001$	r = 0.24; $P < 0.001$	r = 0.24; $P < 0.001$
Daily dairy Ca intake (mg/d)	_	<u>_</u>	r = -0.06; $P = 0.06$	r = -0.12; $P < 0.001$	r = -0.01; $P = 0.73$	r = -0.06; $P = 0.05$
HADS-anxiety score	_	_	_	r = 0.52; $P < 0.001$	r = 0.35; $P < 0.001$	r = 0.34; $P < 0.001$
HADS-depression score	_	_	_	, <u> </u>	r = 0.24; $P < 0.001$	r = 0.23; $P < 0.001$
MSP intensity during the previous	-	-	_	_	-	r = 0.66; $P < 0.001$
month (0–10)						

ISI, Insomnia Severity Index; HADS, Hospital Anxiety and Depression Scale; MSP, musculoskeletal pain.

^{*}One-way ANOVA test, Kruskal-Wallis one-way ANOVA or χ^2 test as appropriate (P-values < 0.05 were considered statistically significant). Data are expressed as frequency (%), mean ± sp or median (25th-75th percentiles).

[†]Post-hoc analysis revealed significant difference between no clinically significant insomnia and subthreshold insomnia.

[‡]Post-hoc analysis revealed significant difference between no clinically significant insomnia and moderate to severe insomnia.

[§]Post-hoc analysis revealed significant difference between subthreshold insomnia and moderate to severe insomnia.

^{*}Pearson's or Spearman's correlation test was used as appropriate (P-values were two-tailed and considered statistically significant at < 0.05).

Table 3 Predictors of ISI score*

Variable	R^2	ANOVA	Model	В	β	P*
101	0.00	E 04.07 D .0.004	Constant	0.72	_	0.62
ISI score 0.22 $F = 31.67$	F = 31.67, P < 0.001	College	-0.03	-0.02	0.52	
			Smoking	1.45	0.09	0.001
			History of medical illness during previous month	0.03	<0.01	0.95
			HADS-anxiety score	0.44	0.35	<0.001
			HADS-depression score	0.12	0.08	0.02
			Daily dairy Ca intake	<-0.01	-0.06	0.03
			Complaint of MSP in any site of the body	-1.22	– 0⋅11	0.04
			MSP intensity during the previous month	0.26	0.12	0.02
			Number of painful body sites	0.32	0.10	0.02

ISI, Insomnia Severity Index; F^{ρ} , squared coefficient of determination; B, unstandardised coefficient; β , standardised coefficient; F, F-statistic; HADS, Hospital Anxiety and Depression Score; MSP, musculoskeletal pain.

Table 4 Predictors of HADS-anxiety and depression scores

Variable	R^2	ANOVA	Model	В	β	<i>P</i> *
HADS-anxiety score	0.45	F=90·15. P <0·001	Constant	-0.80	_	0.35
HADS-anxiety score	0.45	F = 90·15, F < 0·001	Academic level	-0.68	-0.08	0.001
			Gender	1.18	0.13	<0.001
			BMI	0.06	0.06	0.02
			ISI score	0.22	0.27	<0.001
			Daily dairy Ca intake	<0.001	0.03	0.20
			HADS-depression score	0.50	0.43	<0.001
			Complaint of MSP in any site of the body	0.65	0.07	0.10
			MSP intensity during the previous month	0.11	0.06	0.17
			Number of painful body sites	0.18	0.07	0.05
HADS-depression score	0.33	F = 86.79, P < 0.001	Constant	3.54	_	<0.001
·			ISI score	0.05	0.07	0.02
			Daily dairy Ca intake	<-0.01	-0.08	<0.01
			HADS-anxiety score	0.43	0.50	<0.001
			Complaint of MSP in any site of the body	0.06	0.01	0.88
			MSP intensity during the previous month	0.02	0.02	0.75
			Number of painful body sites	0.04	0.02	0.62
			History of medical illness during last month	0.52	0.04	0.10

HADS, Hospital Anxiety and Depression Score; ISI, Insomnia Severity Index; R^2 , squared coefficient of determination; B, unstandardised coefficient; β , standardised coefficient; F, F-statistic; MSP, musculoskeletal pain.

and HADS-depression scores (r=0·27, P-value < 0·001) as well as between ISI and both MSP intensity during the previous month and number of painful body sites (r=0·24, P-values < 0·001). In contrast, daily dairy Ca intake was weakly inversely correlated with both ISI score (r=-0·10, P-value <0·001) and HADS-depression score (r=-0·12, P-value <0·001). HADS-depression score was also moderately correlated with MSP intensity during the previous month (r=0·24, P-value <0·001) and with the number of painful body sites (r=0·23, P-value <0·001). HADS-anxiety score was also moderately correlated with MSP intensity during the previous month (r=0·35, P-value <0·001) and with the number of painful body sites (r=0·34, P-value <0·001).

Predictors of insomnia

Multiple linear regression analysis (Table 3) indicated significant associations between ISI score and smoking (P-value = 0.001), HADS-anxiety score (P-value < 0.001),

HADS-depression score (P-value = 0·02), complaints of MSP (P-value = 0·04), MSP intensity during the previous month (P-value = 0·02) and the number of painful body sites (P-value = 0·02). In contrast, ISI score was significantly inversely associated with daily dairy Ca intake (P-value = 0·03).

Predictors of psychological symptoms (anxiety and depression)

Multiple linear regression analysis (Table 4) indicated that HADS-anxiety score was significantly inversely associated with participants' academic level (P = 0.001) and positively associated with female gender (P-value < 0.001), BMI (P-value = 0.02), ISI score (P-value < 0.001) and HADS-depression score (P-value < 0.001). HADS-depression score was also significantly associated with ISI score (P-value = 0.02) and HADS-anxiety score (P-value < 0.001) and was inversely associated with daily dairy Ca intake (P-value < 0.01).



^{*}Multiple linear regression analysis (P-values < 0.05 were considered statistically significant).

^{*}Multiple linear regression analysis (P-values < 0.05 were considered statistically significant).



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Table 5 Predictors of MSP

Variable	Value	В	SE	OR	CI	P*
Constant		-3.02	0.48	_	_	<0.01
Academic level	Junior students (reference) Senior students	0.11	0.14	1.12	0.85, 1.47	0.41
Gender	Male (reference) Female	0.33	0.15	1.39	1.03, 1.88	0.03
BMI	_	0.04	0.02	1.04	1.01, 1.07	0.02
HADS-anxiety score	_	0.13	0.02	1.14	1.10, 1.19	<0.001
HADS-depression score	_	0.03	0.02	1.04	0.99, 1.08	0.13
ISI score	_	0.03	0.01	1.03	1.00, 1.06	<0.05

HADS, Hospital Anxiety and Depression Scale; MSP, musculoskeletal pain; ISI, Insomnia Severity Index; B, coefficient (intercept). *Binary logistic regression (dependent variable: complaint of MSP in any site of the body), P-values < 0.05 were considered statistically significant.

Predictors of musculoskeletal pain

Binary logistic regression analysis (Table 5) revealed that MSP complaints among participants were significantly associated with female gender (OR 1.39, P-value = 0.03), increased BMI (OR 1.04, *P*-value = 0.02), increased HADSanxiety score (OR 1·14, P-value < 0·001) and increased ISI score (OR 1.03, P-value < 0.05).

Discussion

The current study reports a high prevalence of inadequate Ca intake estimated from daily food consumption among young adults of around 21 years of age. Of concern, the median dairy Ca intake in our cohort of university students was < 1/3 of the recommended daily intake for adults (1000 mg/d)⁽²⁵⁾. Low dietary Ca intake in similar age groups has also been reported by other studies using different methods to assess dietary Ca intake^(26,27). The low dietary Ca intake among young adults could be explained by the wide spread consumption of junk food, especially in universities, that does not contain enough Ca⁽²⁸⁾. In the present study, only dairy food contributions to Ca intake were estimated and no consideration was given to Ca in other foods such as green leafy vegetables or legumes with lower Ca content and poor bioavailability. Ca is highly available in dairy products⁽²⁹⁾, and low dietary Ca intake has been linked to deleterious consequences on the skeleton leading to bone diseases such as osteoporosis⁽²⁵⁾. In addition, low Ca intake could adversely affect other extra-skeletal functions since Ca is involved in muscle contraction, neurotransmission, signalling and other functions⁽³⁰⁾. As such, our hypothesis focused on the relationship between low dairy Ca intake and other inter-related health problems that are commonly manifested in university students including insomnia, anxiety, depression and MSP.

Our major finding was the significant inverse association between insomnia (ISI score) and dairy Ca intake (Table 3). This association suggests that insomnia among university students can be predicted from their low intake of dairy Ca. This finding was also supported by the significant difference in dairy Ca intake between participants

with moderate to severe insomnia and participants with no or subthreshold insomnia (Table 1). The relationship between sleep and Ca could partially be explained by the role of brain Ca in converting the amino acid tryptophan to melatonin, which is considered to be a sleep-inducing hormone⁽⁸⁾. In addition, previous studies have indicated that Ca could be involved in regulating both cycles and duration of sleep in mammals⁽¹⁰⁾. Our results support the increasing evidence that suggests a role for dietary Ca in improving sleep quality and reducing insomnia^(7,8).

Our results have also shown that insomnia among university students can be predicted from anxiety and depression scores, MSP and MSP intensity during the previous month, number of painful body sites and smoking (Table 3). These results were supported by the significant higher anxiety and depression scores in participants with moderate to severe insomnia compared to both participants with no clinically significant insomnia and participants with subthreshold insomnia (Table 1). In addition, a higher percentage of participants who were smokers, and participants with MSP, were classified to have moderate to severe insomnia (Table 1). Moreover, the median MSP intensity and number of painful sites were significantly higher in participants with moderate to severe insomnia compared to both participants with no clinically significant insomnia and participants with subthreshold insomnia (Table 1). The relationship between insomnia and the psychological symptoms of anxiety and depression was previously reported in the literature to be bi-directional^(31,32). This was also supported by our findings as insomnia (ISI score) was a significant predictor for both anxiety and depression (Table 4) and vice versa (Table 3). This suggests that insomnia among university students may lead to anxiety and depression which in turn may cause insomnia. Insomnia and MSP were also previously reported to co-occur and each of them may increase the risk of the other (15). Our results also supported this relationship as insomnia was a significant predictor of MSP, while MSP, MSP intensity and number of painful body sites were significant predictors of insomnia (Tables 3 and 5). The last predictor of insomnia in our study was cigarette smoking. It has been reported that sleep disturbances are more common in cigarette smokers compared



with non-smokers⁽³³⁾. This is explained by the stimulant action of nicotine that can increase alertness⁽³⁴⁾, which may result in insomnia.

One of the main objectives of the current study was to find a link between low dietary Ca estimated from dairy food consumption and insomnia, anxiety, depression and MSP, as we believe that low Ca intake may contribute to the development of these conditions. Our results have shown that dairy Ca intake was inversely associated with insomnia and depression scores but not associated with anxiety scores and complaints of MSP (Tables 3, 4 and 5). These results are not completely consistent with our previous study which was conducted on subjects with non-cardiac musculoskeletal chest pain⁽¹⁴⁾. In that study, low dairy Ca intake was associated with anxiety, and anxiety was associated with pain. This inconsistency may be explained by differences in the study design and age of participants⁽¹⁴⁾. Because participants of the current study were younger and apparently healthy, their low dairy Ca intake may not result in musculoskeletal symptoms. Although there was no significant association between dairy Ca intake and anxiety or MSP, our results suggested an indirect relationship between them. The current study results indicate that anxiety may be affected by low dairy Ca intake through the significant association between anxiety and both insomnia and depression scores, which themselves are associated with low dairy Ca intake. Similarly, MSP may be affected by low dairy Ca intake through the significant association between MSP and insomnia scores, the latter of which is associated with low dairy Ca intake. Therefore, our results maintain support for the presence of a relationship between low dairy Ca intake, insomnia, anxiety, depression and MSP in our study population.

Taken together, the current study has exposed a relationship between low dairy Ca intake and medical conditions including insomnia, anxiety, depression and MSP, all of which are commonly present in university students. The main strengths of the current study were its large sample size and the established methods used to measure the variables of interest. All outcomes were determined using well-validated surveys that have been previously documented in the literature (refer to Materials and methods). The study was mainly limited by its cross-sectional design and the dependence on participants' reliable self-reporting. The study is also limited by the incomplete estimate of total dietary Ca intake, which only included dairy food sources and not that of other foods and beverages with lower Ca content and often poor bioavailability. Dairy foods are the most commonly consumed Ca-rich foods by Jordanian students with highly available Ca. These limitations may prevent us from making conclusions that can be applied to the general population. However, we believe these results are reported for the first time and hope they may encourage other researchers to further investigate the potential causal relationship between low dietary Ca intake and insomnia, anxiety, depression and MSP.

If proven true, further studies should be performed to determine if increasing dietary Ca intake, either via food choices or supplementation, is able to resolve some of these health issues among university students in order to increase both their lifestyle and scientific achievements.

Conclusion

Low dietary Ca intake estimated from dairy food consumption was highly prevalent and associated with insomnia and depression among university students from Jordan. Insomnia in our study participants could be predicted from increased anxiety and depression scores, complaint of MSP and smoking. Both anxiety and depression scores could also be predicted from insomnia scores, supporting a bi-directional association for both anxiety and depression with insomnia. Young adults should be advised to increase dietary Ca intake to achieve the recommended daily amount. It is also required to investigate a potential causal relationship between low dietary Ca and both insomnia and its related co-morbidities.

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