



Epidemiology

# Soy intake and vasomotor menopausal symptoms among midlife women: a pooled analysis of five studies from the InterLACE consortium

Yashvee Dunneram<sup>1</sup> · Hsin-Fang Chung<sup>2</sup> · Janet E. Cade<sup>1</sup> · Darren C. Greenwood<sup>1</sup> · Annette J. Dobson<sup>2</sup> · Ellen S. Mitchell<sup>3</sup> · Nancy F. Woods<sup>4</sup> · Eric J. Brunner<sup>5</sup> · Toyoko Yoshizawa<sup>6</sup> · Debra Anderson<sup>7</sup> · Gita D. Mishra<sup>2</sup>

Received: 21 September 2018 / Revised: 3 January 2019 / Accepted: 16 January 2019 / Published online: 4 February 2019  
© Springer Nature Limited 2019

## Abstract

**Background/objectives** Phytoestrogen rich-foods such as soy may be associated with less frequent/severe vasomotor menopausal symptoms (VMS), although evidence is limited. We thus investigated the associations between the consumption of soy products and soy milk and the frequency/severity of VMS.

**Subjects/methods** We pooled data from 19,351 middle-aged women from five observational studies in Australia, UK, USA, and Japan that contribute to the International Collaboration for a Life course Approach to reproductive health and Chronic disease Events (InterLACE). Information on soy consumption, VMS and covariates were collected by self-report. We included 11,006 women who had complete data on soy consumption, VMS and covariates at baseline for the cross-sectional analysis. For the prospective analysis, 4522 women who were free of VMS at baseline and had complete data on VMS at follow-up were considered. Multinomial logistic regression and binary logistic regression models were used.

**Results** No statistically significant evidence of an association was found between soy products (relative risk ratio (RRR): 0.92, 95% CI: 0.76–1.11) or soy milk (RRR: 1.24, 95% CI: 0.93–1.65) and the likelihood of reporting frequent or severe VMS cross-sectionally. Prospective results indicated that frequent consumption of soy products (odds ratio (OR): 0.63, 95% CI: 0.45–0.89) but not soy milk (OR: 1.11, 95% CI: 0.85–1.45) was associated with lower likelihood of reporting subsequent VMS, after adjustment for socio-demographic and reproductive factors.

**Conclusions** These are the first ever findings from pooled observational data of association between consumption of soy products and VMS.

---

**Sources of support:** Commonwealth Scholarship, Australian National Health and Medical Research Council project grant (APP1027196 & APP1121844)

---

**Supplementary information** The online version of this article (<https://doi.org/10.1038/s41430-019-0398-9>) contains supplementary material, which is available to authorised users.

---

✉ Yashvee Dunneram  
fsyd@leeds.ac.uk

<sup>1</sup> Nutritional Epidemiology Group, School of Food Science and Nutrition, University of Leeds, Leeds, UK

<sup>2</sup> School of Public Health, The University of Queensland, Brisbane, Queensland, Australia

<sup>3</sup> Family and Child Nursing, School of Nursing, University of Washington, Seattle, WA, USA

## Introduction

Menopause, a natural event marking the end of the reproductive life of women, is often accompanied by menopausal symptoms. Vasomotor menopausal symptoms (VMS), including hot flushes and night sweats, are the most common symptoms which arise as a consequence of a decline in endogenous oestrogen levels, in particular during the

<sup>4</sup> Biobehavioral Nursing and Health Systems, School of Nursing, University of Washington, Seattle, WA, USA

<sup>5</sup> Department of Epidemiology and Public Health, University College London, London, UK

<sup>6</sup> Department of Women's Health Nursing, Tohoku University Graduate School of Medicine, Sendai, Japan

<sup>7</sup> Menzies Health Institute Queensland, Griffith University, Gold Coast, Queensland, Australia

perimenopausal and early postmenopausal phases [1, 2]. The frequency and severity of VMS usually, decrease over time, but this varies by individual with symptoms subsiding after a year for some or persisting for over 30 years in others [3]. The frequency/severity of VMS have been linked to various chronic diseases, including cardiovascular disease, osteoporosis and cognitive decline [4, 5].

Phytoestrogen rich-foods such as soy have been associated with less frequent and less severe menopausal symptoms, although evidence is limited [6, 7]. Epidemiological studies which investigated the association between soy intake and the frequency/severity of VMS also demonstrated conflicting results [8, 9]. Moreover, according to a review of 43 randomised controlled trials (RCTs) [1], the positive effect of phytoestrogen supplements on the frequency/severity of hot flushes and night sweats in peri- or postmenopausal women is still inconclusive given the small sample size and potential high risk of bias of the included trials. However, the same review suggested that the effect of genistein (a soy derived isoflavone) was promising [1].

While dietary intake of phytoestrogens is usually, in the form of soy bean, soy bean curd, tofu, tempeh, soy milk and other soy products, most studies have investigated the effects of soy supplements and extracts [10–12]. This study thus sought to elucidate the cross-sectional and prospective associations between soy intake and VMS among peri and postmenopausal women across five studies contributing to the International Collaboration for a Life course Approach to reproductive health and Chronic disease Events (InterLACE) consortium.

## Subjects and methods

### Ethical approval

Written consent was obtained from all participants. All the cohort studies included in the InterLACE consortium have been previously granted ethical approval by the respective ethical committees [13].

### Study participants

The InterLACE consortium includes individual data from ten countries. It involves around 230,000 participants from 20 observational studies with data on women's health (12 of which provided longitudinal data). Further detailed information on InterLACE has been published elsewhere [13, 14]. For the current study, five studies that had information on soy intake (the exposure) and hot flushes and/or night sweats (the outcome) were included: Australian Longitudinal Study on Women's Health (ALSWH) [15],

Healthy Ageing of Women Study (HOW)—Australia, Whitehall II study (WHITEHALL)—UK [16], Seattle Midlife Women's Health Study (SMWHS) [17] and Japanese Midlife Women's Health Study (JMWHS) [18] (Supplementary Table 1). For the cross-sectional analysis, data from 11,006 women who reported VMS (either frequency or severity), consumption frequency of soy products and soy milk and had complete information on confounders (listed below) were included in the analysis. The prospective analysis included data from three studies (ALSWH, HOW and WHITEHALL) ( $n = 10,082$ ). Excluding 5560 women who reported VMS at baseline and those with missing data on VMS, menopausal status and use of hormone therapy at follow-up, 4522 women were considered for the prospective analysis (Supplementary Figure 1).

### Main outcome and exposure variables

VMS was defined as the presence of hot flushes and/ or night sweats. Response options for the frequency of hot flushes and night sweats (over the last 12 months) were 'never, rarely, sometimes, and often' in ALSWH. For the other four studies, the severity of VMS over a shorter period was recorded; HOW, WHITEHALL and JMWHS considered the current severity of VMS, while SMWHS considered the severity of VMS in the last 1–3 months. For example, in HOW and JMWHS the response options for the extent of symptoms were 'not at all, a little, quite a bit, and extremely' and for WHITEHALL the response options were 'not at all, a little, somewhat, and a lot'. The degree of severity was harmonised as 'never, mild, moderate and severe' over a shorter period of time. Since the frequency of VMS was assessed in ALSWH and severity in the remaining four studies, results were presented separately. VMS were further coded dichotomously as 'absent' (never and rarely if reporting frequency; never and mild if reporting severity) and 'present' (sometimes and often if reporting frequency; moderate and severe if reporting severity) for the study-specific and prospective analysis.

Soy products such as tofu, soy beans, tempeh, and soy milk were commonly reported in the five studies. The soy products were combined based on their phytoestrogen contents. Thus, tofu, soy beans, tempeh and soy flour having a high phytoestrogen content were grouped under the soy products category, while soy milk was considered separately [19, 20].

In ALSWH, there were ten consumption frequency options: 'never, less than once per month, 1–3 times per month, 1 time per week, 2 times per week, 3–4 times per week, 5–6 times per week, 1 time per day, 2 times per day and 3 or more times per day'. In the WHITEHALL study, nine consumption frequency options were provided; five in SMWHS and four response categories in HOW and

JMWHS. Therefore for this study, studies having more than four categories were collapsed into four frequency categories: 'never/rarely', 'monthly', 'weekly' and 'daily'. They were further coded dichotomously as 'less frequent' (never/rarely and monthly) and 'frequent' (weekly and daily) given the small number of observations for 'weekly' and 'daily' intake for the prospective analysis.

## Covariates

Categorical variables in the InterLACE study were collapsed into the simplest categories possible so as to include data from as many studies as possible [13]. For example, education level was collated into three categories as  $\leq 10$  years, 11–12 years and  $> 12$  years. Smoking status was grouped as never smokers, past smokers and current smokers. Based on gynaecological surgery and menstrual bleeding patterns, menopausal status was collated into five categories to include (1) hysterectomy/oophorectomy, (2) unknown due to hormone use (menopausal hormone therapy or oral contraceptive hormones before reaching menopause), (3) premenopause (regular menstruation in the last 3 and 12 months), (4) perimenopause (menses in the past 3 months and changes/irregularity in menstrual patterns in the past 12 months; or no menses in the previous 3 months but menses in the preceding 11 months) and (5) natural postmenopause (amenorrhoea for at least 12 months). Current use of menopausal hormone therapy (e.g., oestrogen) was categorised as yes and no.

## Statistical analysis

As the result of different assessments (frequency or severity) and different recall periods (in the past 12 months or in a more recent period) for VMS, studies were grouped as: (1) frequency of VMS in the past 12 months (ALSWH) and (2) severity of VMS over a shorter time period (HOW, WHITEHALL, SMWHS and JMWHS). The associations between soy consumption and VMS were first examined separately for the two different designs, followed by the overall estimates.

Multinomial logistic regression models with four categories of outcome for VMS (never, rarely/mild, sometimes/moderate and often/severe) were used to investigate the cross-sectional associations between frequency of consumption of soy products and soy milk with frequency/severity of VMS at baseline. The VMS category 'never' was used as the reference group for the outcome, and the soy consumption category 'never' was used as the reference group for the exposure. Relative risk ratios (RRR) and 95% confidence intervals (CI) were estimated. According to the minimally sufficient set of adjustments, smoking status, education level, menopausal status and race/ethnicity were

identified as confounders using a directed acyclic graph (DAG) (Supplementary Figure 2) and were adjusted for in the regression models. However, race/ethnicity was not included in the model as participants from ALSWH (96.5%), HOW (95.1%), WHITEHALL (88.1%) and SMWHS (88.1%) were mainly Caucasians, and in JMWHS all the participants were Japanese. Concurrent menopausal hormone therapy use was included in the model given its potential effect on the frequency/severity of VMS [21]. The models were thus adjusted for menopausal status and concurrent menopausal hormone therapy use (model 1) and additionally adjusted for other potential covariates including education level and smoking status (model 2). 'Study' was included as a fixed effect to account for differences in levels of VMS between studies and as a stratification variable to account for correlation of individuals within studies.

Due to small numbers of participants in the four categories of exposure and outcome in individual studies, dichotomised soy consumption (frequent and less frequent) and dichotomised VMS (presence and absence) were used for the study-specific and prospective analyses. To examine between-study heterogeneity in the effect size estimates, study-specific logistic regression and random-effects meta-analysis were used with the estimates adjusted for all the covariates in model 2.

For the prospective analysis based on three studies (ALSWH, HOW and WHITEHALL), logistic regression models with the binary outcome for VMS (presence and absence) were fitted, adjusted for all the covariates in model 2. In addition, a sensitivity analysis was conducted to investigate the association between soy consumption and subsequent risk of VMS at follow-up with all the women included ( $n = 10,082$ ), but adjusting for their baseline VMS, given that a large proportion of women were excluded in the prospective analysis due to the presence of VMS at baseline. Analyses were performed using STATA 14 (StataCorp LP, College Station, TX). All statistical tests were two sided.

## Results

A total of 11,006 women reported their consumption frequency of soy and VMS, and also had complete data on the covariates. The median age of the women at baseline was 52 years (interquartile range: 51–54) (Supplementary Table 1). Table 1 shows the baseline characteristics of the participants in each study. The majority of the participants were Caucasians-Australians/New-Zealanders (57.5%), had 10 years or less of education (46.3%), and never smoked (60.9%). Nearly, 30% of the women were naturally postmenopausal, and 26.5% were currently using menopausal hormone therapy. Across HOW, WHITEHALL, SMWHS

**Table 1** Baseline characteristics of participants

Characteristics	Overall	ALSWH	HOW	WHITEHALL	SMWHS	JMWHS
<i>n</i>	11,006	7373	563	2146	174	750
<i>Race/ethnicity</i>						
Caucasian-Australian/ New Zealand	6323 (57.5)	5853 (79.4)	470 (83.5)	0 (0.0)	0 (0.0)	0 (0.0)
Caucasian-European	3163 (28.7)	1207 (16.4)	65 (11.6)	1891 (88.1)	0 (0.0)	0 (0.0)
Caucasian-American/ Canadian	202 (1.8)	54 (0.7)	0 (0.0)	0 (0.0)	148 (85.1)	0 (0.0)
Japanese	756 (6.9)	6 (0.1)	0 (0.0)	0 (0.0)	0 (0.0)	750 (100.0)
Chinese and other Asians	166 (1.5)	144 (2.0)	7 (1.2)	0 (0.0)	15 (8.6)	0 (0.0)
Others	396 (3.6)	109 (1.5)	21 (3.7)	255 (11.9)	11 (6.3)	0 (0.0)
<i>Birth year (n = 11,002)</i>						
<1940	856 (7.8)	N/A	N/A	855 (39.8)	1 (0.6)	N/A
1940–1949	7354 (66.8)	5410 (73.4)	475 (85.0)	1034 (48.2)	81 (46.6)	354 (47.2)
≥1950	2792 (25.4)	1963 (26.6)	84 (15.0)	257 (12.0)	92 (52.9)	396 (52.8)
<i>Education level</i>						
≤10 years	5096 (46.3)	3568 (48.4)	286 (50.8)	1171 (54.6)	0 (0.0)	71 (9.5)
11–12 years	2177 (19.8)	1269 (17.2)	92 (16.3)	345 (16.1)	24 (13.8)	447 (59.6)
>12 years	3733 (33.9)	2536 (34.4)	185 (32.9)	630 (29.4)	150 (86.2)	232 (30.9)
<i>Marital status (n = 10,225)</i>						
Married	7927 (77.5)	6028 (82.0)	427 (76.3)	1357 (63.3)	115 (66.1)	N/A
Separated/divorced/ widowed	1597 (15.6)	1099 (15.0)	106 (18.9)	340 (15.9)	52 (29.9)	N/A
Single	701 (6.9)	221 (3.0)	27 (4.8)	446 (20.8)	7 (4.0)	N/A
<i>Body mass index (n = 10,425)</i>						
Normal weight (<25 kg/m <sup>2</sup> )	5071 (48.6)	3048 (43.9)	233 (43.3)	1068 (52.7)	90 (51.7)	632 (85.8)
Overweight (25–29.9 kg/m <sup>2</sup> )	3259 (31.3)	2297 (33.1)	165 (30.7)	654 (32.3)	47 (27.0)	96 (13.0)
Obese (≥30 kg/m <sup>2</sup> )	2095 (20.1)	1604 (23.1)	140 (26.0)	305 (15.1)	37 (21.3)	9 (1.2)
<i>Smoking status</i>						
Never	6707 (60.9)	4505 (61.1)	356 (63.2)	1108 (51.6)	89 (51.2)	649 (86.5)
Past smoker	2704 (24.6)	1782 (24.2)	158 (28.1)	667 (31.1)	67 (38.5)	30 (4.0)
Current smoker	1595 (14.5)	1086 (14.7)	49 (8.7)	371 (17.3)	18 (10.3)	71 (9.5)
<i>Menopausal status</i>						
Hysterectomy/ oophorectomy	2598 (23.6)	2001 (27.1)	165 (29.3)	344 (16.0)	6 (3.5)	82 (10.9)
Unknown due to hormone use	1721 (15.6)	1346 (18.3)	46 (8.2)	265 (12.4)	47 (27.0)	17 (2.3)
Premenopause	1315 (12.0)	636 (8.6)	22 (3.9)	463 (21.6)	44 (25.3)	150 (20.0)
Perimenopause	2090 (19.0)	1484 (20.1)	76 (13.5)	390 (18.2)	53 (30.5)	87 (11.6)
Natural menopause	3282 (29.8)	1906 (25.9)	254 (45.1)	684 (31.9)	24 (13.8)	414 (55.2)
<i>Current use of menopausal hormone therapy</i>						
No	8085 (73.5)	5043 (68.4)	369 (65.5)	1813 (84.5)	134 (77.0)	726 (96.8)
Yes	2921 (26.5)	2330 (31.6)	194 (34.5)	333 (15.5)	40 (23.0)	24 (3.2)
<i>Frequency or severity of hot flushes</i>						
Never	4443 (40.4)	2249 (30.5)	323 (57.4)	1344 (62.6)	118 (67.8)	409 (54.5)
Rarely or mild	2009 (18.3)	1183 (16.1)	160 (28.4)	388 (18.1)	29 (16.7)	249 (33.2)
	2608 (23.7)	2241 (30.4)	59 (10.5)	233 (10.9)	15 (8.6)	60 (8.0)

**Table 1** (continued)

Characteristics	Overall	ALSWH	HOW	WHITEHALL	SMWHS	JMWHS
<i>n</i>	11,006	7373	563	2146	174	750
Sometimes or moderate						
Often or severe	1946 (17.7)	1700 (23.1)	21 (3.73)	181 (8.4)	12 (6.9)	32 (4.3)
<i>Frequency or severity of night sweats</i>						
Never	5510 (50.1)	2996 (40.6)	358 (63.6)	1458 (67.9)	137 (78.7)	561 (74.8)
Rarely or mild	1813 (16.5)	1157 (15.7)	136 (24.2)	339 (15.8)	22 (12.6)	159 (21.2)
Sometimes or moderate	2183 (19.8)	1914 (26.0)	52 (9.2)	190 (8.9)	5 (2.9)	22 (2.9)
Often or severe	1500 (13.6)	1306 (17.7)	17 (3.0)	159 (7.4)	10 (5.8)	8 (1.1)
<i>Frequency or severity of vasomotor symptoms<sup>a</sup></i>						
Never	4049 (36.8)	2034 (27.6)	285 (50.6)	1251 (58.3)	112 (64.4)	367 (48.9)
Rarely or mild	2099 (19.1)	1212 (16.4)	184 (32.7)	388 (18.1)	31 (17.8)	284 (37.9)
Sometimes or moderate	2728 (24.8)	2312 (31.4)	66 (11.7)	269 (12.5)	15 (8.6)	66 (8.8)
Often or severe	2130 (19.4)	1815 (24.6)	28 (5.0)	238 (11.1)	16 (9.2)	33 (4.4)
<i>Consumption frequency of soy products</i>						
Never/rarely	9239 (84.0)	6590 (89.4)	475 (84.4)	2047 (95.4)	127 (73.0)	0 (0.0)
Monthly	491 (4.5)	357 (4.8)	50 (8.9)	62 (2.9)	0 (0.0)	22 (2.9)
Weekly	820 (7.5)	357 (4.8)	34 (6.0)	35 (1.6)	36 (20.7)	358 (47.7)
Daily	456 (4.1)	69 (0.9)	4 (0.7)	2 (0.1)	11 (6.3)	370 (49.3)
<i>Consumption frequency of soy milk (n = 10,954)</i>						
Never/rarely	9860 (90.0)	6634 (90.0)	460 (85.2)	2103 (98.2)	147 (84.5)	516 (70.8)
Monthly	156 (1.4)	29 (0.4)	21 (3.9)	17 (0.8)	0 (0.0)	89 (12.2)
Weekly	237 (2.2)	110 (1.5)	11 (2.0)	19 (0.9)	13 (7.5)	84 (11.5)
Daily	701 (6.4)	596 (8.1)	48 (8.9)	3 (0.1)	14 (8.1)	40 (5.5)

Data are presented as *n* (%); N/A—not applicable

<sup>a</sup>Vasomotor menopausal symptoms were defined as having hot flashes, night sweats, or both

and JMWHS which measured the severity of VMS, WHITEHALL had the highest percentage of women who reported 'severe' VMS (11.1%), while JMWHS (Japanese) had the lowest percentage (4.4%). In the ALSWH study, 24.6% reported 'often' for the frequency of VMS. In this predominantly Caucasian population, 80–90% of the women reported that they never consumed soy products or soy milk. Across the individual studies, JMWHS had the largest percentage of women who reported 'daily' and 'weekly' soy product consumption (49.3% and 47.7%, respectively) (Table 1). Comparing baseline characteristics of women included in the prospective analysis and those excluded due to loss to follow-up, the excluded women were less educated and more likely to be obese and current smokers at baseline. They were more likely to be post-menopausal and less likely to report frequent/severe VMS compared to women with complete follow-up data (Supplementary Table 2).

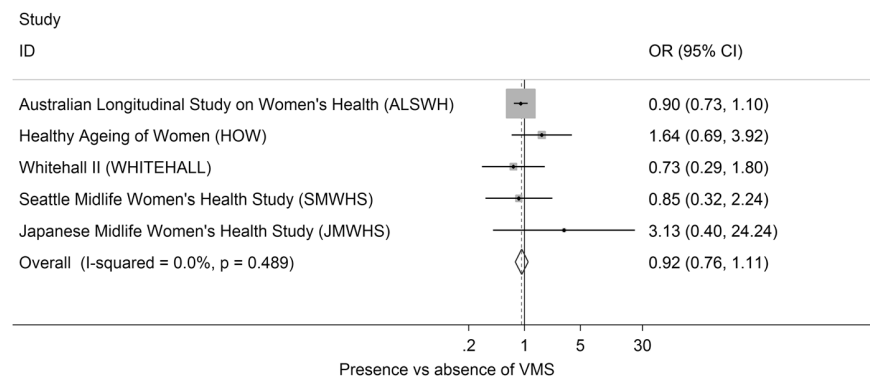
For the cross-sectional analysis, women with 'weekly' and 'daily' consumption of soy products were less likely to

report frequent/severe VMS compared to those with 'never/rarely' consumption (11.7% vs. 20.5% and 6.4% vs. 20.5%, respectively) (Table 2). However, after adjusting for covariates and study differences, no clear evidence of an association was found between soy product consumption and the degree of VMS. Similarly, there was no clear evidence of an association observed for ALSWH or the other four studies. For soy milk consumption, women with a daily consumption were more likely to report frequent/severe VMS compared to women who reported 'never/rarely' consumption (RRR: 1.56, 95% CI: 1.24–1.96). A similar pattern for 'daily' consumption and risk of frequent/severe VMS was observed in ALSWH (RRR: 1.39, 95% CI: 1.10–1.77) and the other four studies (RRR: 3.09, 95% CI: 1.47–6.50).

When using dichotomised exposure and outcome variables for the study-specific analysis, the pooled estimate of association between frequent soy product consumption and the presence of VMS was 0.92 (95% CI: 0.76–1.11), with no statistically significant heterogeneity between studies,

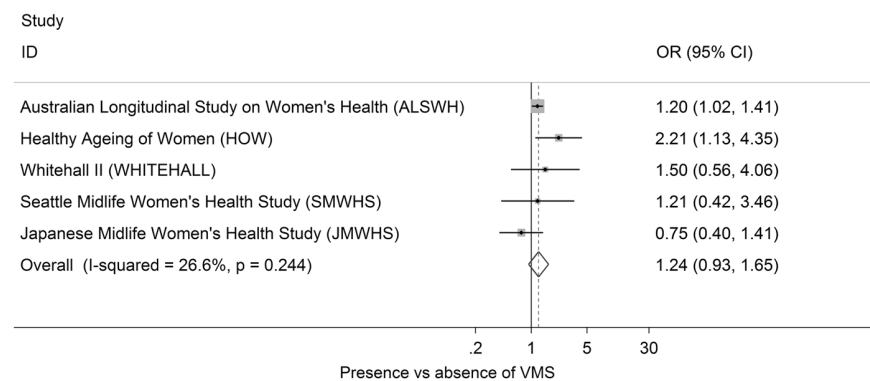
**Table 2** Cross-sectional association of soy products and soy milk consumption frequency with the frequency/severity of vasomotor menopausal symptoms at baseline

		VMS (hot flushes and night sweats) (%)				Model 1 <sup>a</sup> RRR (95% CI)				Model 2 <sup>b</sup> RRR (95% CI)			
<i>n</i>	Never	Rarely/mild	Sometimes/moderate	Often/severe	Never	Rarely/mild	Sometimes/moderate	Often/severe	Rarely/mild	Often/severe	Sometimes/moderate	Often/severe	
<b>Soy consumption frequency</b>													
<b>Soy products</b>													
<i>ALSWH<sup>c</sup> (n = 7373)</i>													
Never/rarely	6590	27.3	16.6	31.4	24.8	Reference	Reference	Reference	Reference	Reference	Reference	Reference	
Monthly	357	29.1	14.9	29.4	26.6	Reference 0.81 (0.58–1.14)	Reference 0.84 (0.63–1.11)	Reference 0.96 (0.71–1.29)	Reference 0.80 (0.57–1.13)	Reference 1.03 (0.76–1.38)	Reference 0.85 (0.64–1.13)	Reference 1.03 (0.76–1.38)	
Weekly	357	31.9	14.6	33.1	20.5	Reference 0.72 (0.52–1.02)	Reference 0.85 (0.65–1.11)	Reference 0.66 (0.49–0.90)	Reference 0.71 (0.51–1.00)	Reference 0.73 (0.53–0.99)	Reference 0.86 (0.66–1.13)	Reference 0.73 (0.53–0.99)	
Daily	69	29.0	20.3	33.3	17.4	Reference 1.13 (0.57–2.26)	Reference 0.97 (0.52–1.79)	Reference 0.64 (0.31–1.33)	Reference 1.11 (0.56–2.21)	Reference 0.69 (0.33–1.43)	Reference 0.98 (0.53–1.81)	Reference 0.69 (0.33–1.43)	
<i>HOW, WHITEHALL, SMWHS, JMWHS<sup>c</sup> (n = 3633)</i>													
Never/rarely	2649	57.3	20.7	12.1	9.9	Reference	Reference	Reference	Reference	Reference	Reference	Reference	
Monthly	134	53.0	24.6	11.9	10.5	Reference 0.97 (0.63–1.51)	Reference 1.08 (0.60–1.93)	Reference 1.45 (0.76–2.70)	Reference 1.02 (0.65–1.59)	Reference 1.70 (0.90–3.21)	Reference 1.23 (0.68–2.21)	Reference 1.70 (0.90–3.21)	
Weekly	463	52.3	31.3	11.5	5.0	Reference 1.01 (0.65–1.56)	Reference 1.22 (0.71–2.09)	Reference 0.86 (0.42–1.75)	Reference 1.05 (0.68–1.63)	Reference 0.95 (0.46–1.95)	Reference 1.35 (0.79–2.31)	Reference 0.95 (0.46–1.95)	
Daily	387	47.3	41.6	6.7	4.4	Reference 1.33 (0.81–2.19)	Reference 0.77 (0.38–1.55)	Reference 0.86 (0.37–2.02)	Reference 1.41 (0.86–2.32)	Reference 1.00 (0.43–2.34)	Reference 0.89 (0.44–1.80)	Reference 1.00 (0.43–2.34)	
<i>OVERALL (n = 11,006)</i>													
Never/rarely	9239	35.9	17.8	25.8	20.5	Reference	Reference	Reference	Reference	Reference	Reference	Reference	
Monthly	491	35.6	17.5	24.6	22.2	Reference 0.85 (0.65–1.12)	Reference 0.89 (0.69–1.14)	Reference 1.03 (0.79–1.35)	Reference 0.87 (0.66–1.13)	Reference 1.14 (0.88–1.49)	Reference 0.93 (0.72–1.20)	Reference 1.14 (0.88–1.49)	
Weekly	820	43.4	24.0	20.9	11.7	Reference 0.81 (0.63–1.05)	Reference 0.95 (0.75–1.20)	Reference 0.70 (0.53–0.93)	Reference 0.85 (0.66–1.09)	Reference 0.82 (0.62–1.09)	Reference 1.03 (0.81–1.30)	Reference 0.82 (0.62–1.09)	
Daily	456	44.5	38.4	10.8	6.4	Reference 1.11 (0.79–1.56)	Reference 0.77 (0.51–1.16)	Reference 0.65 (0.40–1.05)	Reference 1.18 (0.85–1.65)	Reference 0.78 (0.49–1.24)	Reference 0.85 (0.57–1.28)	Reference 0.78 (0.49–1.24)	
<b>Soy milk</b>													
<i>ALSWH (n = 7369)</i>													
Never/rarely	6634	27.9	16.5	31.3	24.3	Reference	Reference	Reference	Reference	Reference	Reference	Reference	
Monthly	29	34.5	13.8	41.4	10.3	Reference 0.68 (0.21–2.17)	Reference 1.05 (0.45–2.48)	Reference 0.34 (0.09–1.25)	Reference 0.69 (0.21–2.20)	Reference 0.37 (0.10–1.38)	Reference 1.08 (0.46–2.54)	Reference 0.37 (0.10–1.38)	
Weekly	110	22.7	19.1	35.5	22.7	Reference 1.41 (0.79–2.54)	Reference 1.38 (0.83–2.32)	Reference 1.17 (0.66–2.08)	Reference 1.42 (0.79–2.56)	Reference 1.27 (0.71–2.26)	Reference 1.41 (0.84–2.36)	Reference 1.27 (0.71–2.26)	
Daily	596	24.7	15.3	30.7	29.4	Reference 1.02 (0.78–1.34)	Reference 1.08 (0.86–1.36)	Reference 1.33 (1.05–1.68)	Reference 1.03 (0.78–1.35)	Reference 1.39 (1.10–1.77)	Reference 1.09 (0.87–1.37)	Reference 1.39 (1.10–1.77)	



**Fig. 1** Forest plot of study-specific effect estimates of the cross-sectional association between consumption frequency of soy products and the presence of vasomotor menopausal symptoms at baseline. Soy product consumption was coded dichotomously as 'frequent' (weekly and daily) and 'less frequent' (never/rarely and monthly) and vasomotor symptoms as 'present' (sometimes and often if reporting frequency; moderate and severe if reporting severity) and 'absent' (never

and rarely if reporting frequency; never and mild if reporting severity) given the small number of observations in each study. Odds ratios (ORs) are presented on a log scale. Effect estimates were adjusted for menopausal status, current use of menopausal hormone therapy, education level, and smoking status. VMS – vasomotor menopausal symptoms



**Fig. 2** Forest plot of study-specific effect estimates of the cross-sectional association between consumption frequency of soy milk and the presence of vasomotor menopausal symptoms at baseline. Soy milk consumption was coded dichotomously as 'frequent' (weekly and daily) and 'less frequent' (never/rarely and monthly) and vasomotor symptoms as 'present' (sometimes and often if reporting frequency; moderate and severe if reporting severity) and 'absent' (never and

rarely if reporting frequency; never and mild if reporting severity) given the small number of observations in each study. Odds ratios (ORs) are presented on a log scale. Effect estimates were adjusted for menopausal status, current use of menopausal hormone therapy, education level and smoking status. VMS – vasomotor menopausal symptoms

test for heterogeneity:  $P = 0.49$ ,  $I^2 = 0\%$  (Fig. 1). For the association between frequent consumption of soy milk and the presence of VMS, the pooled OR estimate was 1.24 (95% CI: 0.93–1.65) with no statistically significant heterogeneity between the studies (test for heterogeneity:  $P = 0.24$ ,  $I^2 = 26.6\%$ ) (Fig. 2).

For the prospective analysis, the overall estimates suggest that women who had frequent soy product consumption were less likely to report the incidence of VMS at follow-up (OR: 0.63, 95% CI: 0.45–0.89) (Table 3). A consistent pattern was observed in ALSWH (OR: 0.63, 95% CI: 0.44–0.90) and the other four studies (OR: 0.60, 95% CI: 0.18–1.97). There was no clear evidence of an

association between frequent consumption of soy milk and incident VMS at follow-up (OR: 1.11, 95% CI: 0.85–1.45). The sensitivity analysis with all the women included demonstrated a similar or weaker association between soy consumption and subsequent VMS, even after adjusting for baseline VMS (Table 4).

## Discussion

This pooled study demonstrated no clear evidence of an association between consumption frequency of soy products and VMS in the cross-sectional analysis. However, in the

**Table 3** Prospective association of soy product and soy milk consumption frequency with the presence of vasomotor menopausal symptoms at the follow-up survey

Soy consumption	<i>n</i>	VMS <sup>a</sup> (%)	Crude OR (95% CI)	Model 1 <sup>b</sup> OR (95% CI)	Model 2 <sup>c</sup> OR (95% CI)
<b>Soy products</b>					
<i>ALSWH (n = 2,852)</i>					
Less frequent <sup>d</sup>	2688	35.5	Reference	Reference	Reference
Frequent	164	26.2	0.65 (0.45–0.92)	0.63 (0.44–0.91)	0.63 (0.44–0.90)
<i>HOW, WHITEHALL (n = 1670)</i>					
Less frequent	1625	12.4	Reference	Reference	Reference
Frequent	45	6.7	0.56 (0.17–1.85)	0.58 (0.18–1.91)	0.60 (0.18–1.97)
<i>OVERALL (n = 4522)</i>					
Less frequent	4313	26.8	Reference	Reference	Reference
Frequent	209	22.0	0.64 (0.45–0.90)	0.63 (0.45–1.88)	0.63 (0.45–0.89)
<b>Soy milk</b>					
<i>ALSWH (n = 2849)</i>					
Less frequent	2608	34.9	Reference	Reference	Reference
Frequent	241	35.7	1.04 (0.79–1.37)	1.05 (0.79–1.38)	1.04 (0.79–1.38)
<i>HOW, WHITEHALL (n = 1655)</i>					
Less frequent	1614	12.2	Reference	Reference	Reference
Frequent	41	17.1	2.01 (0.85–4.78)	2.08 (0.86–4.99)	2.18 (0.91–5.23)
<i>OVERALL (n = 4504)</i>					
Less frequent	4222	26.2	Reference	Reference	Reference
Frequent	282	33.0	1.09 (0.84–1.43)	1.10 (0.84–1.43)	1.11 (0.85–1.45)

Logistic regression models were used to estimate odds ratios (OR) and 95% confidence intervals (95% CI).

<sup>a</sup>VMS was defined as ‘presence of VMS’ for ‘never’ and ‘rarely/mild’ VMS and ‘absence of VMS’ for ‘sometimes/moderate’ and ‘often/severe’ VMS

<sup>b</sup>Model 1 was adjusted for menopausal status and current use of menopausal hormone therapy at follow-up

<sup>c</sup>Model 2 was adjusted for model 1 along with other covariates including smoking status, and education level

<sup>d</sup>Soy consumption frequency was defined as ‘frequent’ for ‘daily’ and ‘weekly’ consumption and ‘less frequent’ for ‘monthly’ and ‘never/rarely’ consumption

prospective analysis, women with frequent consumption of soy products were less likely to report subsequent VMS. Furthermore, there was no evidence of an association between consumption of soy milk and frequency/severity of VMS both cross-sectionally (Figs. 1 and 2) and prospectively (Table 3).

Our prospective analysis showed an association between frequent consumption of soy products and decreased odds of VMS at follow-up, though this was attenuated when baseline VMS was taken into account. Similarly, a Japanese community-based study in which women were followed for 6 years found that soy products intake alleviated hot flushes [9]. Several RCTs have investigated the association between some type of substance containing dietary soy (e.g., soy extract in capsule or tablet form, soy powder or soy protein added to diets) and its effect on hot flushes. While some demonstrated a reduction in the frequency/severity of hot flushes [10, 22–24], others have shown contradictory findings [25, 26]. According to a review study, the dose of genistein, in particular, was associated with a reduction of

the symptoms rather than total isoflavone [27]. The oestrogen-like properties of soy food due to the isoflavones content have been linked to the protective effect on VMS. A decrease in the number of ovarian follicles and consequent fall in oestrogen level could be the underlying hormonal aetiology of VMS [28, 29]. However, the effect of phytoestrogens in reducing VMS remains unclear [30]. One of the possible mechanism of action is the structural similarity of isoflavones to that of oestradiol could confer oestrogenic or anti-oestrogenic effects depending on the circulating oestrogen level by binding to oestrogen receptors [31, 32]. The relative decline in oestrogen level leads to higher circulating norepinephrine levels and an upregulation of serotonin receptors which mediate hot flushes in menopausal women. By binding to oestrogen receptors, isoflavones help to restore the oestrogen level, and causes subsequent changes in norepinephrine and serotonin levels, thus reducing the propensity of hot flushes [33].

Our pooled data did not show a clear association between soy milk consumption and frequency/severity of VMS. The



**Table 4** Sensitivity analysis for the prospective association between soy consumption and likelihood of reporting vasomotor symptoms at follow-up

Soy consumption	<i>n</i>	VMS <sup>a</sup> (%)	Crude OR (95% CI)	Model 1 <sup>b</sup> OR (95% CI)	Model 2 <sup>c</sup> OR (95% CI)	Model 3 <sup>d</sup> OR (95% CI)
<b>Soy products</b>						
<i>ALSWH (n = 6603)</i>						
Less frequent <sup>e</sup>	6235	54.8	Reference	Reference	Reference	Reference
Frequent	368	48.6	0.78 (0.63–0.96)	0.76 (0.62–0.94)	0.79 (0.64–0.98)	0.79 (0.63–1.00)
<i>HOW, WHITEHALL (n = 2251)</i>						
Less frequent	2194	21.4	Reference	Reference	Reference	Reference
Frequent	57	19.3	1.05 (0.54–2.07)	0.99 (0.50–1.97)	1.05 (0.53–2.08)	1.04 (0.54–1.99)
<i>OVERALL (n = 8854)</i>						
Less frequent	8429	46.1	Reference	Reference	Reference	Reference
Frequent	425	44.7	0.80 (0.65–0.98)	0.78 (0.63–0.96)	0.81 (0.66–1.00)	0.81 (0.65–1.01)
<b>Soy milk</b>						
<i>ALSWH (n = 6599)</i>						
Less frequent	5970	54.1	Reference	Reference	Reference	Reference
Frequent	629	57.6	1.15 (0.97–1.36)	1.15 (0.97–1.36)	1.17 (0.99–1.38)	1.06 (0.89–1.27)
<i>HOW, WHITEHALL (n = 2233)</i>						
Less frequent	2175	21.2	Reference	Reference	Reference	Reference
Frequent	58	31.0	2.63 (1.43–4.84)	2.44 (1.29–4.60)	2.65 (1.40–5.00)	2.10 (1.07–4.13)
<i>OVERALL (n = 8832)</i>						
Less frequent	8145	45.3	Reference	Reference	Reference	Reference
Frequent	687	55.3	1.21 (1.03–1.43)	1.20 (1.01–1.41)	1.22 (1.04–1.44)	1.10 (0.92–1.32)

Logistic regression models were used to estimate odds ratios (OR) and 95% confidence intervals (95% CI).

<sup>a</sup>VMS was defined as ‘presence of VMS’ for ‘never’ and ‘rarely/mild’ VMS and ‘absence of VMS’ for ‘sometimes/moderate’ and ‘often/severe’ VMS

<sup>b</sup>Model 1 was adjusted for menopausal status and current use of menopausal hormone therapy at follow-up

<sup>c</sup>Model 2 was adjusted for model 1 along with other covariates including smoking status and education level

<sup>d</sup>Model 3 was adjusted for model 2 along with baseline VMS

<sup>e</sup>Soy consumption frequency was defined as ‘frequent’ for ‘daily’ and ‘weekly’ consumption and ‘less frequent’ for ‘monthly’ and ‘never/rarely’ consumption

source of dietary isoflavones may also contribute to the observed effect since processing methods tend to alter the phytoestrogen contents of soy products [34]. For instance, the total isoflavone content in soy beans (103 mg per 100 g), tempeh (18 mg per 100 g) and tofu (27 mg per 100 g) is much higher than that in soy milk (3 mg per 100 g) [20]. The overall low-consumption frequency of soy milk among the participants and its low isoflavone content could possibly explain this finding.

The main drawback of our study is the variation in assessments used by the different studies. Soy consumption was measured as frequency, with no information on quantities. Moreover, for the consumption of soy milk, the cross-sectional nature of some of the studies and lack of evidence of a significant association from the prospective analysis, mean that we cannot confirm a temporal relationship between soy milk consumption and VMS. There also might be possibility of residual confounding, e.g., by factors not measured in the studies. One weakness of data harmonisation is the collapsing of the variables of interest into the simplest level of detail in order to incorporate information from as many studies as possible, leading to loss of

statistical power as well as potential misclassification of the degree of VMS and frequency of soy consumption. For instance, studies like ALSWH and WHITEHALL had ten and nine frequency options respectively for consumption of soy that were collapsed to four categories for this analysis. In addition, the frequency of VMS was reported in ALSWH over a longer period of time (12 months), and the other four studies recorded the severity of VMS over a shorter period that limited our ability to pool data. Despite these limitations the pooled results showed considerable homogeneity as shown in the forest plots and the low values for the statistic  $I^2$ .

Furthermore, our study had several strengths that ranged from the inclusion of a large number of women across different geographic regions and cultures that allowed greater generalisability of the results. This is also, to our knowledge, the first pooled study consisting of women’s health studies from four different countries examining an association between soy products and soy milk with frequency/severity of VMS. We also included women who had a hysterectomy, oophorectomy, and/or were currently using hormones that could provide a better estimate of the

prevalence of VMS. In addition, the individual data available in the InterLACE enabled harmonisation of the variables of interest using common definitions, coding and cut points not normally possible with meta-analyses of published results. Harmonisation of the data further reduces the between-study heterogeneity. A consistent approach to confounder adjustment was used for the regression models along with careful selection of the confounders using a DAG, thus reducing the probability of the results being affected by uncontrolled confounders.

While menopause is an inevitable phenomenon in a woman's life cycle, the frequency and severity of VMS show marked variations [35]. VMS are reported by around 75% of postmenopausal women globally, with a minority reporting severe symptoms [36, 37]. Findings from this study provide some evidence that frequent consumption of soy products (e.g., soy beans, tofu and tempeh) as part of the usual diet may be associated with a reduced risk of subsequent VMS. However, frequent consumption of soy milk did not appear to be associated with subsequent VMS. As justified by potential mechanisms in previous studies, our findings could prompt RCTs testing the effects of dietary soy intake in particular on VMS as opposed to earlier RCTs which have mainly considered the effects of soy extracts and supplements.

**Acknowledgements** The data on which this research are based were drawn from five observational studies. The research included data from the ALSWH, the University of Newcastle, Australia, and the University of Queensland, Australia. We are grateful to the Australian Government Department of Health for funding and to the women who provided the survey data. SMWHS was supported in part by grants from the National Institute of Nursing Research. HOW and JMWHS (also called Australian and Japanese Midlife Women's Health Study) were supported by the Queensland University of Technology Early Career Research Grant and the JSPS Grant-in-aid for Scientific Research. The Whitehall II study is supported by grants from the Medical Research Council (K013351), British Heart Foundation (BHF RG/16/11/32334) and US National Institutes on Aging (R01AG013196 and R01AG034454). All study teams would like to thank the participants for volunteering their time to be involved in the respective studies. The findings and views in this paper are not those from the original studies or their respective funding agencies.

**Funding** InterLACE project is funded by the Australian National Health and Medical Research Council project grant (APP1027196). GDM is supported by the Australian National Health and Medical Research Council Principal Research Fellowship (APP1121844). YD is in receipt of a scholarship from the Commonwealth Scholarship Commission, UK. The funders had no role in study design, data collection, and analysis, decision to publish, or preparation of the manuscript.

**Author contributions** The authors' responsibilities were as follows—GDM: conceived the study; YD, HFC and GDM: designed the research and had primary responsibility for the final content; JEC, DCG, AJD, ESM, NFW, EJB, TY and DA: contributed to the data; YD: performed the statistical analysis and wrote the manuscript; HFC, DCG, JEC, AJD and GDM: provided statistical input, helped with

interpretation of the results and reviewed the manuscript for important intellectual content; and all authors: read and approved the final manuscript.

## Compliance with ethical standards

**Conflict of interest** JEC is the director of a university spin out company, Dietary Assessment Ltd. The remaining authors declare that they have no conflict of interest.

**Publisher's note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

## References

- Lethaby A, Marjoribanks J, Kronenberg F, Roberts H, Eden J, Brown J. Phytoestrogens for menopausal vasomotor symptoms. *Cochrane Database Syst Rev*. 2013;Cd001395.
- Gold EB, Colvin A, Avis N, Bromberger J, Greendale GA, Powell L, et al. Longitudinal analysis of the association between vasomotor symptoms and race/ethnicity across the menopausal transition: study of women's health across the nation. *Am J Public Health*. 2006;96:1226–35.
- Rapkin AJ. Vasomotor symptoms in menopause: physiologic condition and central nervous system approaches to treatment. *Am J Obstet Gynecol*. 2007;196:97–106.
- Pines A. Vasomotor symptoms and cardiovascular disease risk. *Climacteric*. 2011;14:535–6.
- Biglia N, Cagnacci A, Gambacciani M, Lello S, Maffei S, Nappi RE. Vasomotor symptoms in menopause: a biomarker of cardiovascular disease risk and other chronic diseases? *Climacteric*. 2017;20:306–12.
- Levis S, Griebeler ML. The role of soy foods in the treatment of menopausal symptoms. *J Nutr*. 2010;140:2318s–2321s.
- Baber R. Phytoestrogens and post reproductive health. *Maturitas*. 2010;66:344–9.
- Sievert LL, Morrison L, Brown DE, Reza AM. Vasomotor symptoms among Japanese-American and European-American women living in Hilo, Hawaii. *Menopause*. 2007;14:261–9.
- Nagata C, Takatsuka N, Kawakami N, Shimizu H. Soy product intake and hot flashes in Japanese women: results from a community-based prospective study. *Am J Epidemiol*. 2001;153:790–3.
- Carmignani LO, Pedro AO, Costa-Paiva LH, Pinto-Neto AM. The effect of dietary soy supplementation compared to estrogen and placebo on menopausal symptoms: a randomized controlled trial. *Maturitas*. 2010;67:262–9.
- Khaothiar L, Ricciotti HA, Li L, Pan W, Schickel M, Zhou J, et al. Daidzein-rich isoflavone aglycones are potentially effective in reducing hot flashes in menopausal women. *Menopause*. 2008;15:125–32.
- Ye YB, Wang ZL, Zhuo SY, Lu W, Liao HF, Verbruggen M, et al. Soy germ isoflavones improve menopausal symptoms but have no effect on blood lipids in early postmenopausal Chinese women: a randomized placebo-controlled trial. *Menopause*. 2012;19:791–8.
- Mishra GD, Chung HF, Pandeya N, Dobson AJ, Jones L, Avis NE, et al. The InterLACE study: design, data harmonization and characteristics across 20 studies on women's health. *Maturitas*. 2016;92:176–85.
- Mishra GD, Anderson D, Schoenaker DA, Adami HO, Avis NE, Brown D, et al. InterLACE: A New International Collaboration for a Life Course Approach to Women's Reproductive Health and Chronic Disease Events. *Maturitas*. 2013;74:235–40.

15. Lee C, Dobson AJ, Brown WJ, Bryson L, Byles J, Warner-Smith P, et al. Cohort Profile: the Australian Longitudinal Study on Women's Health. *Int J Epidemiol.* 2005;34:987–91.
16. Marmot M, Brunner E. Cohort Profile: the Whitehall II study. *Int J Epidemiol.* 2005;34:251–6.
17. Mitchell ES, Woods NF. Cognitive symptoms during the menopausal transition and early postmenopause. *Climacteric.* 2011;14:252–61.
18. Anderson D, Yoshizawa T, Gollschewski S, Atogami F, Courtney M. Menopause in Australia and Japan: effects of country of residence on menopausal status and menopausal symptoms. *Climacteric.* 2004;7:165–74.
19. Nutrient Data L. USDA database for the isoflavone content of selected foods, release 2.0. Beltsville, Md: U.S. Department of Agriculture; 2008.
20. Thompson LU, Boucher BA, Liu Z, Cotterchio M, Kreiger N. Phytoestrogen content of foods consumed in Canada, including isoflavones, lignans, and coumestrol. *Nutr Cancer.* 2006;54:184–201.
21. Stuenkel CA, Davis SR, Gompel A, Lumsden MA, Murad MH, Pinkerton JV, et al. Treatment of symptoms of the menopause: an Endocrine Society Clinical Practice Guideline. *J Clin Endocrinol Metab.* 2015;100:3975–4011.
22. Welty FK, Lee KS, Lew NS, Nasca M, Zhou JR. The association between soy nut consumption and decreased menopausal symptoms. *J Women's Health.* 2007;16:361–9.
23. Cheng G, Wilczek B, Warner M, Gustafsson JA, Landgren BM. Isoflavone treatment for acute menopausal symptoms. *Menopause.* 2007;14:468–73.
24. Nahas EA, Nahas-Neto J, Orsatti FL, Carvalho EP, Oliveira ML, Dias R. Efficacy and safety of a soy isoflavone extract in postmenopausal women: a randomized, double-blind, and placebo-controlled study. *Maturitas.* 2007;58:249–58.
25. Burke GL, Legault C, Anthony M, Bland DR, Morgan TM, Naughton MJ, et al. Soy protein and isoflavone effects on vasomotor symptoms in peri- and postmenopausal women: the Soy Estrogen Alternative Study. *Menopause.* 2003;10:147–53.
26. Penotti M, Fabio E, Modena AB, Rinaldi M, Omodei U, Viganó P. Effect of soy-derived isoflavones on hot flushes, endometrial thickness, and the pulsatility index of the uterine and cerebral arteries. *Fertil Steril.* 2003;79:1112–7.
27. Williamson-Hughes PS, Flickinger BD, Messina MJ, Empie MW. Isoflavone supplements containing predominantly genistein reduce hot flash symptoms: a critical review of published studies. *Menopause.* 2006;13:831–9.
28. Sturdee DW. The menopausal hot flush—anything new? *Maturitas.* 2008;60:42–49.
29. Pachman DR, Jones JM, Loprinzi CL. Management of menopause-associated vasomotor symptoms: current treatment options, challenges and future directions. *Int J Women's Health.* 2010;2:123–35.
30. Moreira AC, Silva AM, Santos MS, Sardão VA. Phytoestrogens as alternative hormone replacement therapy in menopause: what is real, what is unknown. *J Steroid Biochem Mol Biol.* 2014;143:61–71.
31. Rietjens I, Louise J, Beekmann K. The potential health effects of dietary phytoestrogens. *Br J Pharmacol.* 2017;174:1263–80.
32. Kuiper GG, Lemmen JG, Carlsson B, Corton JC, Safe SH, van der Saag PT, et al. Interaction of estrogenic chemicals and phytoestrogens with estrogen receptor beta. *Endocrinology.* 1998;139:4252–63.
33. Morrow PKH, Mattair DN, Hortobagyi GN. Hot flashes: a review of pathophysiology and treatment modalities. *Oncologist.* 2011;16:1658–64.
34. Setchell KDR, Cole SJ. Variations in isoflavone levels in soy foods and soy protein isolates and issues related to isoflavone databases and food labeling. *J Agric Food Chem.* 2003;51:4146–55.
35. Luoto R. Hot flushes and quality of life during menopause. *BMC Women's Health.* 2009;9:13.
36. Monteleone P, Mascagni G, Giannini A, Genazzani AR, Simoncini T. Symptoms of menopause—global prevalence, physiology and implications. *Nat Rev Endocrinol.* 2018;14:199–215.
37. Woods NF, Mitchell ES. Symptoms during the perimenopause: prevalence, severity, trajectory, and significance in women's lives. *Am J Med.* 2005;118(Suppl 12B):14–24.