

Sleep and Methylation of Estrogen Receptor Genes, *ESR1* and *GPER*, in Healthy Middle-Aged and Older Women: Findings from the Women 40+ Healthy Aging Study

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Purpose: Sleep problems in middle-aged and older women are very common and have been associated with menopause-related changes in estrogen levels. However, not all women experience sleep problems as they enter perimenopause, and epigenetic mechanisms might contribute to the differences in sleep quality within this population. In this study, we hypothesized that increased methylation of two estrogen receptor (ER) genes (*ESR1* and *GPER*) would be associated with increased sleep problems in healthy pre-, peri-, and postmenopausal women, either directly or indirectly through the experience of vasomotor symptoms (VMS).

Materials and Methods: In 130 healthy women aged 40–73 years, we assessed DNA methylation from dried blood spots (DBS). Women rated their sleep quality using the Pittsburgh Sleep Quality Index (PSQI), and VMS using the Menopause Rating Scale (MRS).

Results: Higher percentage methylation of *ESR1* was associated with increased sleep problems, mediated by VMS, even after controlling for age, menopausal status, body mass index, estradiol levels, depressive symptoms, and caffeine consumption. There was no significant association between *GPER* methylation and either sleep problems or VMS.

Conclusion: The study findings support an association between increased *ESR1* methylation and sleep problems through increased VMS among healthy women aged 40–73 years.

Keywords: sleep problems, VMS, *ESR1*, *GPER*, DNA methylation, healthy middle-aged and older women

Introduction

Sleep is an important determinant of quality of life during aging, even among healthy individuals.^{1,2} Sleep problems in the form of insomnia, difficulty falling asleep, lower sleep efficiency, or frequent and early awakening become increasingly common with advancing age, and affect as many as 40–56% of middle-aged and older women.³ Menopausal status (MS) is strongly associated with sleep problems, with peri- and postmenopausal women reporting more sleep problems than premenopausal women.⁴ Changes in levels of ovarian hormones such as estrogens are important contributors to the occurrence and severity of sleep problems as women enter perimenopause.⁵ Indeed, besides their role in women's reproductive functions, estrogens exert a vast range of biological effects. In the central nervous system, estrogens contribute to the control of circadian rhythms and body temperature,^{5,6}

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and their declining and fluctuating levels have been repeatedly associated with sleep problems and vasomotor symptoms (VMS).^{5–10} VMS, such as hot flashes and night sweats, are primarily thermoregulatory phenomena and are the main cause of sleep problems.^{3,11} Although VMS are a hallmark of perimenopause, they can persist for several years after the last menstrual period. Therefore, they may continue to affect sleep even in advanced age.¹² However, while all aging women experience major hormonal changes, not all women experience increased sleep problems.¹³ This suggests that genetic and environmental factors may play a role in the exacerbation of sleep problems among middle-aged and older women.^{14–18}

Epigenetic modifications are increasingly recognized as potential mechanisms underlying the etiology and phenotypic variation of multiple diseases.¹⁹ It has also been suggested that epigenetic modifications may constitute underlying mechanisms of the gene-environment interplay affecting sleep and menopause-related symptoms.^{14,20,21} DNA methylation is a key epigenetic mechanism, which regulates gene expression through DNA modifications at cytosines in cytosine-guanine dinucleotides (CpGs).²² Increased DNA methylation of regulatory regions of estrogen receptor (ER) genes has been shown to decrease levels of ERs.^{6,23,24} ERs mediate the effects of estrogens in the body, including the estrogenic contributions to the control of circadian rhythms and body temperature.^{25–27} While the ER α and the ER β mainly mediate the genomic effects of estrogens, the more recently discovered *G protein*-coupled estrogen receptor 1 (GPER) mediates rapid non-genomic estrogen effects.²⁸ Several lines of evidence from animal studies indicate that decreased levels of the three ERs may contribute to increase sleep problems and VMS.^{29–31}

To the best of our knowledge, no study has yet investigated the associations between ER gene methylation and sleep problems in middle-aged and older women. DNA methylation levels of specific regions of ER genes may contribute to alter sleep quality among women as they reach midlife. Therefore, in this study, we hypothesized that increased levels of ER gene methylation would be associated with increased sleep problems in healthy pre-, peri-, and postmenopausal women, either directly, or indirectly through VMS. We focused on the ER α gene (*ESR1*) and GPER gene (*GPER*) based on knowledge of the involvement of both genomic and non-genomic estrogen effects in sleep problems and VMS, and given the presumed greater role of ER α compared to ER β in controlling circadian rhythms and body temperature in women.^{29,32}

Materials and Methods

Inclusion and Exclusion Criteria

Women who took part in this study were recruited in the context of a larger investigation, the “Women 40+ Healthy Aging Study”.^{33–36} Women were recruited using flyers, social media (eg: Facebook), and journal articles. The prospective participants completed an online self-screening to verify their eligibility for the study. Women were included in the study if they reported being free of any acute or chronic somatic disease or mental disorder, and if they had not received any psychotherapeutic or psychopharmacological treatment during the last 6 months. In addition, women had to report good, very good, or excellent subjective health status. Further exclusion criteria were a pregnancy in the last 6 months, menopause due to surgical removal of the ovaries or the uterus, premature menopause, current use of oral contraceptives or hormone therapy. After completion of the online self-screening, eligible women were reached telephonically by a member of the study team. The telephone call aimed, first, to ensure that only women suitable for the study were included, and second, to answer open questions about the study procedure. Participants were invited to a laboratory session during which saliva and blood samples were collected. All sessions started at 7:45 a.m. and followed a standardized protocol.

Participants

In total, 130 healthy Caucasian women aged 40–73 years took part in the study. The women lived in Switzerland, and the majority were Swiss or from a German-speaking neighboring country. Most of the women were married or living together with their partner (68.5%), and had vocational education or a college/university degree (79.2%). Participants were divided into three subgroups with respect to their MS according to the Stages of Reproductive Aging Workshop +10 (STRAW) criteria:³⁷ 1) premenopausal (39.2%, n= 51), if the menstrual cycle was regular, 2) perimenopausal (12.3%, n= 16), if the cycle length was variable, with a variability among cycles of at least 7 days, or if the interval between cycles was > 60 days, and 3) postmenopausal (48.5%, n= 63), if no bleeding had occurred in at least the last 12 months. Women using hormone therapies for reducing menopausal symptoms were excluded during the online self-screening. After enrolment, women were screened for the use of other medications. Five women reported taking botanical therapies such as Cimifemin[®] (2), Cimifemin[®]

forte (1), Premens (1), and a plant-based preparation (no pharmaceutical name provided; 1). In addition, one woman reported taking Eltroxin (hypothyroidism) and one woman reported taking Tamoxifen (breast cancer prevention). The results were not affected by the exclusion of these participants from the statistical analyses, and the sample of this study includes participants reporting the use of the above-mentioned medications. After receiving written and oral information about the study, all participants signed written informed consent forms. The study (BASEC Nr 2016–01591) was evaluated by the cantonal ethics committee of the Canton of Zürich (KEK Zürich, Zürich, Switzerland). Approval from the cantonal ethics committee was not required because the assessment of relationships between biological and physiological parameters in healthy aging individuals was not considered to fall within the scope of the “Humanforschungsgesetz” (HFG) (Art. 2 HFG and Art. 3 HFG).

Sleep Problems

Sleep problems were assessed using the German version of the Pittsburgh Sleep Quality Index (PSQI).³⁸ The scale assesses sleep problems over a one-month time interval and comprises 19 items combined into seven component scores: perceived sleep quality (“how would you rate your sleep quality overall?”), sleep latency (eg: “cannot get to sleep within 30 minutes”), sleep duration (“how many hours of actual sleep do you get at night?”), sleep efficiency (number of hours slept/number of hours spent in bed), sleep disturbances (eg: “wake up in the middle of the night or early morning”, “had bad dreams”), use of sleep medication (eg: “how often have you taken medicines to help you sleep?”), and daytime dysfunction (eg: “how much of a problem has it been for you to keep up enough enthusiasm to get things done?”). Each component is weighted equally from 0 to 3, and the seven component scores are summed up to obtain a total score ranging from 0 to 21. A score above 5 is considered a marker of sleep problems. The single total score is frequently used in the literature, and acceptable reliability and validity in different populations have been reported.³⁹ However, the model including only perceived sleep quality, sleep efficiency and daytime dysfunction (3-factor model) seems to better reflect good versus poor sleep quality in healthy peri- and early postmenopausal women with hot flashes, and in older adults.^{39,40} Therefore, in the statistical analyses of the present study, we used both the total score and the 3-factor model of the PSQI.

Descriptive statistics of sleep problems in the overall sample and based on menopausal groups are reported in Table 1.

Vasomotor Symptoms

VMS were measured using one item (“hot flashes/sweating”) of the somato-vegetative subscale of the German version of the Menopause Rating Scale (MRS).⁴¹ VMS are rated on a 5-point Likert scale, ranging from “0” (no symptoms) to “4” (very severe symptoms). The prevalence of VMS was assessed as follows: 0 (no symptoms), 1–4 (symptoms). Descriptive statistics of VMS for all women and separately for menopausal groups are reported in Table 1.

Blood Sampling

Blood samples were collected at 8:00 am during the laboratory session at the University of Zurich under standardized conditions. For pre- and perimenopausal women, sampling was conducted in the early follicular phase. Blood was collected from fingertips and spotted onto S&S 903 Whatman® paper cards (GE Healthcare, Little Chalfont, Buckinghamshire, United Kingdom), following the manufacturer’s instructions. Blood spots were dried at room temperature for approximately 3 h. Dried blood spots (DBS) were stored at –20°C until biochemical analysis.

Table 1 Descriptive Statistics

N	All	PRE	PERI	POST
	130	51	16	63
Sleep problems (mean/SD)	4.8/2.5	4.2/2.4	5.8/2.6	5/2.5
VMS (mean/SD)	1.8/1	1.3/0.5	2.4/1.1	2/1.2
ESR1 methylation (%; mean/SD)	76.7/12	79/8.4	78.1/7.9	74.5/14.7
GPER methylation (%; mean/SD)	15.3/7	14.8/1.8	19.2/3.8	14.8/1.8
Age (mean/range)	53.2/40-73	45.1/40-57	51.4/47-56	60.2/50-73
BMI (kg/m ² ; mean/SD)	23/3.6	22.8/3.9	21.5/3.4	23.7/3.3
E2 (nmol/L; mean/SD)	5.6/4.5	8.2/4.7	5.7/4.5	3.7/3.2
Depressive symptoms (mean/SD)	8.3/5.9	7.4/5.5	10.1/7.3	8.6/5.7
Caffeine (cups/day; mean/SD)	2.5/2	2.2/0.2	2.1/0.4	2.7/0.1

Note: The percentages of *ESR1* and *GPER* methylation correspond to the mean of methylation percentages of the 9 CpGs and 22 CpGs assessed in the *ESR1* and *GPER* sequences, respectively.

Abbreviations: PRE, premenopausal, PERI, perimenopausal, POST, postmenopausal, VMS, vasomotor symptoms, *ESR1*, estrogen receptor 1 gene, *GPER*, G protein-coupled estrogen receptor gene, BMI, body mass index, E2, estradiol, SD, standard deviation.

DNA Methylation

Genomic DNA was extracted from DBS, using the QIAamp DNA Investigator Kit (QIAGEN, Hilden, Germany), according to the manufacturer's instructions, and eluted in a final volume of 30 μ L of RNase-free water. We obtained a total DNA yield ranging from 41 to 168 ng, as measured with Qubit (Thermo Fischer Scientific, Waltham, MA, USA). The DBS technology was reported to provide highly reliable results in the context of methylation studies.⁴² Genomic DNA (41–168 ng) was bisulfite-converted using the EZ 96-DNA Methylation-Gold kit (Zymo Research, Irvine, CA, USA) and eluted in 20 μ L of water. A first amplification was performed on the bisulfite-converted DNA using the Kapa HiFi Uracil+ master mix and primers that were designed manually or using the MethPrimer software.⁴³ Primers (Table 2) included the universal primer sequences CS1/CS2 at the 5' ends of the forward and reverse primers, and were designed in order to target a specific sequence of the *ESR1* shore of promoter C (hg 38; chr6:151805523–151805822, Figure 1A) and the *GPER* promoter (hg 38; chr7:1087059–1087533, Figure 1B). DNA methylation in these regions was previously associated with altered gene expression and function.^{44,45} The PCR was performed with the following parameters: 95°C for 3 min [98°C for 20 sec – annealing temperature for 15 sec – 72°C for 15 sec] x 40 cycles, 72°C for 40 sec. In order to certify that primers were specifically amplifying bisulfite-converted DNA, genomic DNA was included in the PCR, and complete conversion was confirmed by including wild primers. Next, the PCR amplicons were purified using the E-gels 2% size selection (Thermo Fisher Scientific, Waltham, MA, USA). Purified amplicons were indexed with a single barcode (Fluidigm, San Francisco, California, USA) using a second PCR of 10 cycles (annealing temperature, 60°C). Indexed amplicons were pooled and again purified. The purified library was then quantified using the Agilent 2200 Tape Station instrument, with HS DNA 1000 reagents (Agilent Scientific Instruments,

Santa Clara, CA, USA), and finally diluted to the concentration of 2nM. The final library was denatured in NaOH, mixed with 12% PhiX, diluted to 12 pM in HT1 buffer, and finally sequenced on an Illumina MiSeq sequencer using the v3 kit (300 base pairs paired-end).

As described previously, adaptors and primers were trimmed from the reads using the default settings of Trimmomatic v0.35 (licensed under GPL V3 and available at <http://www.usadellab.org/cms/index.php?page=trimmomatic>), and low-quality reads defined by a Phred quality score <30 were also discarded.^{47,48} Using Bismark program (v0.19.0), the remaining sequencing reads were aligned to the target regions and the number of unconverted and converted cytosine was extracted for each interrogated CpG. After summing up counts of unconverted and converted cytosine for each CpG, we removed all samples showing a coverage (=total count) lower than 100x.⁴⁷ For *ESR1*, six samples did not reach the 100x threshold and were therefore excluded. For the remaining samples, coverage ranged from 157 to 42'709 (Mean= 6'5601, SD= 5'968). The coverage of most samples for *GPER* was less than 100x, and only 34 samples could be retained in the analyses. For these 34 samples, coverage ranged from 350 to 39'942 (Mean= 10'647, SD= 9553). The methylation percentage for each CpG was calculated as the unconverted (cytosine) read count divided by the total read count (cytosine + thymine).

The *ESR1* shore and *GPER* methylation data are openly available in “Dryad” at <https://doi.org/10.5061/dryad.51c59zw51>.

One CpG cluster was identified for *ESR1*, and therefore the mean of the nine assessed CpGs was used in the statistical analyses. For *GPER*, two main clusters were identified. However, the results were similar when using the two clusters or the mean of all 22 CpGs assessed for *GPER*. Therefore, only results obtained using the mean level across all 22 CpGs are presented in this study. Descriptive statistics of *ESR1* and *GPER* methylation

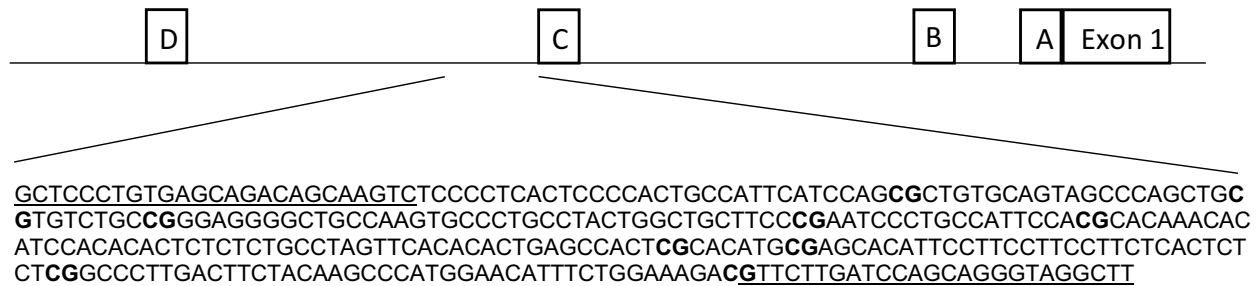
Table 2 PCR Primers Used for Amplification of DNA Sequences in the *ESR1* and *GPER*

Gene	Forward Primer	Reverse Primer	GRCh38	Ta (°C)
<i>ESR1</i>	ACACTGACGACATGGTTCTACA NNN GTTTTTGTGAGTAGATAGTAAGTT	TACGGTAGCAGAGACTTGGTCT NNN AAACCTACCCTACTAAATCAAAAAC	chr6:151805523–151805822	58
<i>GPER</i>	ACACTGACGACATGGTTCTACA NNNAGTGGAAATTTAAATGGTTAGTA	TACGGTAGCAGAGACTTGGTCT NNNACAATCCAAACAATTCAAAATTTATTT	chr7:1087059–1087533	60

Note: Universal primer CS1= ACACTGACGACATGGTTCTACA, universal primer CS2= TACGGTAGCAGAGACTTGGTCT.

Abbreviations: PCR, polymerase chain reaction; *ESR1*, estrogen receptor alpha gene; *GPER*, G protein-coupled estrogen receptor gene; GRCh38, Genome Reference Consortium Human Build 38 Organism; Ta, annealing temperature.

A



B

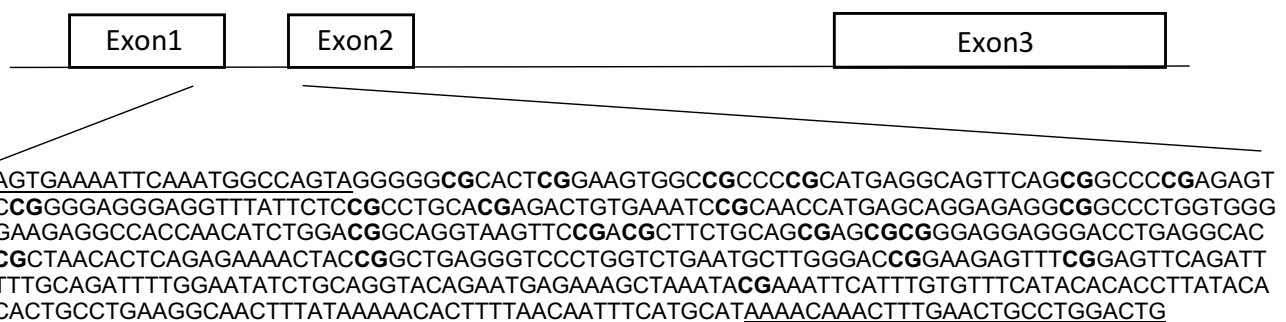


Figure 1 Schematic figures of *ESRI* (A) and *GPER* (B) promoter regions. (A) The DNA sequence of *ESRI*, chr6: 151805523–151805822, is located in a CpG island shore of promoter C, and it has been described as an enhancer (ID GH06|151804) of promoter A.⁴⁴ This region has been reported in humans and rats by Champagne et al, 2006,²³ Gardini et al, 2020,³⁵ lanov et al, 2017,⁴⁶ and Tsuboi et al, 2017.⁴⁴ (B) The DNA sequence of *GPER*, chr7:1087059–1087533 is located in a CpG island across exons 1 and 2. This region has been assessed in Weissenborn et al, 2017.⁴⁵ White boxes represent exons or promoter regions. Underlined sequences correspond to the primers position. Bold “CG” correspond to the Interrogated CpGs.

levels are reported in Table 1 for the entire sample and the three menopausal stages, respectively.

Potential Covariates and Confounders

In the statistical analyses we adjusted for age, MS, body mass index (BMI), levels of estradiol (E2), depressive symptoms, and caffeine consumption.^{8,9,49-57} Depressive symptoms were assessed using the German version of the Center for Epidemiological Studies Depression scale (CES-D), which is a commonly used self-report measure of depressed symptoms during the past one to 2 weeks.⁵⁸ It consists of 20 items (eg, “I felt hopeful about the future”, “I felt that people disliked me”), rated on a 4-point Likert scale ranging from 0 (rarely or none of the time [less than 1 day]) to 3 (most or all

of the time [five to 7 days]). The total score can range from 0 to 60, with higher scores indicating higher depressive symptoms. A cut-off score of 16 or above is often used as a marker of risk for clinical depression.⁵⁸ Levels of E2 concentrations were measured in saliva using an enzyme-linked immunosorbent assay (IBL international, Hamburg, Germany), following the manufacturer’s instructions. Saliva samples were collected at 8:00 am into 2-mL SaliCaps (IBL International GmbH, Hamburg, Germany) immediately before the collection of blood samples. Intra- and inter-assay coefficients were below 13.3% and 14.8%, respectively, and the assay’s analytical sensitivity was 1.1 pmol/L. Values greater than three standard deviations around the mean were considered as outliers (n= 4) and therefore

removed from the analyses. Salivary E2 strongly correlates with free serum E2, which is the portion available for estrogenic effects.^{59,60} BMI was calculated by dividing the weight in kilograms by the height in meters squared, and caffeine consumption was determined as the number of cups of coffee the women drink per day.

Procedure and Statistical Analyses

Descriptive statistics were calculated to assess the prevalence and severity of sleep problems and VMS among the three menopausal groups (pre-, peri and postmenopausal women). We used Chi-squared test in order to test the prevalence of sleep problems and VMS, and one-way analysis of variance (ANOVA) to test their severity. In the Chi-square test post-hoc analysis, p -values were estimated from unstandardized Z values, according to García-perez et al (2003).⁶¹ A p -value below 0.05 was considered statistically significant.

In order to assess the effects of *ESR1*/*GPER* methylation on sleep problems, considering VMS as potential mediators, we conducted simple mediation analyses using the PROCESS macro (model 4)⁶² in SPSS. The mediation model allows the examination of the independent contribution of direct and indirect effects through three paths (a_i , b_i , c' , see Figure 2): Path a_i represents the association between the predictor (X) and the mediator (M_i), path b_i represents the association between M_i and the outcome variable (Y), and path c' represents the direct effect of X on Y , after adjusting for all other variables included in the model. Finally, the indirect effect of X on Y through M_i is defined as $a_i \times b_i$. Direct and indirect effects were assessed using bootstrapping, with the b coefficient providing an index of the magnitude of the effect. Bootstrapping generated a confidence interval (CI). When the 95% asymmetric CI did not include the value 0, direct and indirect effects were considered significant. Bootstrapping has the advantages of not requiring normal distribution of data and of a low vulnerability to type 1 error.⁶² In a second step, MS was dummy-coded, with the premenopausal group (compared to peri- and postmenopausal groups) and perimenopausal and postmenopausal groups (compared to the premenopausal group) as dummies, to investigate whether effects of the

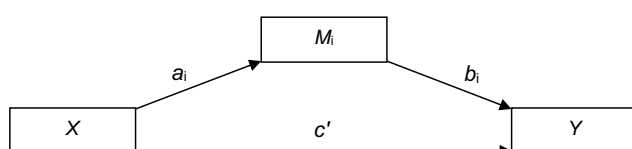


Figure 2 Path diagram (Macro Model 4) according to Hayes (2017).⁶²

mediation analysis differed between premenopausal women and the group including peri- and postmenopausal women. All analyses were conducted using SPSS (IBM Statistics, version 24.0, Armonk, NY: IBM Corp.).

Results

Levels of mean methylation across the investigated CpGs of *ESR1* and *GPER* for the three menopausal groups are provided in Table 1. In addition, all variables assessed in this study are provided in this table.

Descriptive Statistics of Sleep Problems Among Menopausal Groups

Almost one third of the sample (30.5%) experienced sleep problems (score of PSQI > 5) during the previous month. Sleep problems were reported by 24% (n= 12) of premenopausal women, 31.3% (n= 5) of perimenopausal women, and 35.5% (n= 22) of postmenopausal women. The difference in the prevalence of sleep problems among the three menopausal groups was not statistically significant [χ^2 (2, N= 128)= 1.73, p = 0.421]. However, there was a trend toward significance regarding the difference in the severity of sleep problems among the groups [F(2125)= 2.94, p = 0.056], with greater sleep problems in peri- (p = 0.028) and postmenopausal women (p = 0.087) compared to premenopausal women.

Descriptive Statistics of Vasomotor Symptoms Among Menopausal Groups

VMS were reported by 48.1% of all women. VMS were experienced by 27% (n= 14) of premenopausal women, 81.3% (n= 13) of perimenopausal women, and 56.5% (n= 35) of postmenopausal women. Among the three groups, there was a statistically significant difference in the prevalence of VMS [χ^2 (2, N= 129)= 17.49, p < 0.001]. The number of premenopausal women experiencing VMS was significantly lower than the number of peri- and postmenopausal women experiencing VMS (p < 0.001). The difference in the severity of VMS among menopausal groups was also statistically significant [F (2126)= 10.38, p < 0.001], with peri- (p < 0.001) and postmenopausal women (p < 0.001) reporting greater VMS severity compared to premenopausal women. No significant difference in VMS severity was detected between peri- and postmenopausal women (p = 0.130).

ESR1 Methylation and Sleep Problems

Using the total score of the PSQI as outcome variable, the results indicated that *ESR1* methylation predicted sleep problems indirectly through VMS (b = 0.011, 95% CI [0.0013,

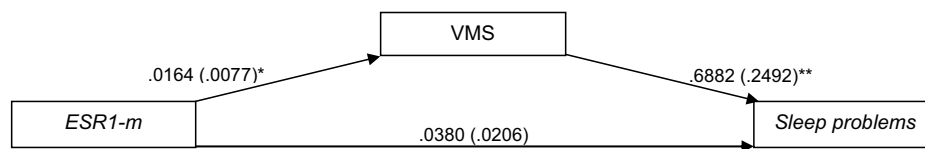


Figure 3 Path diagram of the mediation model (Macro Model 4),⁶² illustrating the associations between *ESR1* methylation (*X*) and sleep disturbances (*Y*) through VMS (*M*). Estradiol (E2), menopausal status (MS), depressive symptoms, body mass index (BMI), age and caffeine consumption were included in the model and controlled in each of the three paths (*a*, *b*, *c*). Values outside parentheses= unstandardized *b* coefficient; values in parentheses= standard error; *= *p*-value < 0.05; **= *p*-value < 0.01.

Abbreviations: *ESR1-m*, estrogen receptor 1 methylation; VMS, vasomotor symptoms.

0.0299]), while the direct effect of *ESR1* methylation on sleep problems was not significant ($b = 0.025$, 95% CI [-0.100, 0.0597], Figure 3). The results of this mediation model are presented in Table 3. The results of the sensitivity analysis indicated that most of the mediation effect came from peri- and postmenopausal women ($b = 0.0127$, 95% CI [0.0005, 0.0246]), compared to premenopausal women ($b = 0.0045$, 95% CI [-0.0020, 0.0121]).

Similar results were obtained when using the 3-factor model. The indirect effect of *ESR1* methylation on the global scores of perceived sleep quality, sleep efficiency and daytime dysfunction through VMS was statistically significant

($b = 0.007$, 95% CI [0.0014, 0.0187]), while the direct effect was not significant ($b = 0.011$, 95% CI [-0.0103, 0.0324]).

GP ER Methylation and Sleep Problems

Using the total score of the PSQI as outcome variable, the results indicated that *GP ER* methylation was not predictive of sleep problems, either directly ($b = 0.073$, $t(23) = 1.08$, 95% CI [-0.0664, 0.2124]), or indirectly through VMS ($b = -0.037$, 95% CI [-1.083, 0.0416], Figure 4). The results of this mediation model are presented in Table 4.

Likewise, when testing the effect of *GP ER* methylation on the 3-factor model of the PSQI, the direct effect of

Table 3 Results of the Mediation Model with *ESR1* Methylation as Predictor, 95% Bias-Corrected Confidence Interval Predicting Sleep Quality

Outcome Variable: VMS	b	se	t	p	LLCI	ULCI
<i>ESR1-m</i> (<i>a</i> ₁)	0.0175	0.0078	2.2550	0.0261	0.0021	0.0329
CV						
Age	-.0516	0.0169	-3.0573	0.0028	-.0850	-.0182
MS	0.7049	0.1640	4.2969	0.0000	0.3798	1.0300
BMI	0.0194	0.0149	1.2970	0.1974	-.0102	0.0490
E2	-.0018	0.0224	-.0790	0.9372	-.0462	0.0427
Depressive symptoms	-.0151	0.0155	-.9704	0.3340	-.0458	0.0157
Caffeine	0.0201	0.0575	0.3502	0.7269	-.0938	0.1340
Outcome variable: sleep problems (PSQI)						
VMS (<i>b</i> ₁)	0.6301	0.2112	2.9837	0.0035	0.2116	1.0487
CV						
Age	0.0586	0.0389	1.5055	0.1351	-.0186	0.1358
MS	-.3619	0.3927	-.9215	0.3588	-1.1401	0.4164
BMI	0.0178	0.0333	0.5331	0.5950	-.0483	0.0838
E2	0.0147	0.0497	0.2962	0.7676	-.0838	0.1132
Depressive symptoms	0.2033	0.0345	5.8904	0.0000	0.1349	0.2717
Caffeine	-.2224	0.1274	-1.7456	0.0837	-.4748	0.0301
Direct/indirect effects on sleep problems (PSQI)						
Direct effect (<i>c'</i>)	0.0249	0.0176	1.4159	0.1597	-.0100	0.0597
Indirect effect (<i>a</i> ₁ <i>x</i> <i>b</i> ₁)	0.0110	0.0073			0.0013	0.0299

Notes: *a*₁ represents the association between *ESR1-m* and VMS; *b*₁ represents the association between VMS and sleep quality; *c'* represents the direct effect of *ESR1-m* on *Y*, after adjusting for all other variables included in the model; *a*₁*x**b*₁ represents the indirect effect of *ESR1-m* on sleep quality through VMS; *b* represents the unstandardized regression coefficient.

Abbreviations: se, standard error; BootLLCI, bootstrapping lower limit confidence interval; BootULCI, bootstrapping upper limit confidence interval; VMS, vasomotor symptoms, *ESR1-m*, estrogen receptor alpha gene methylation; CV, control variables; MS, menopausal status; BMI, body mass index; E2, estradiol.

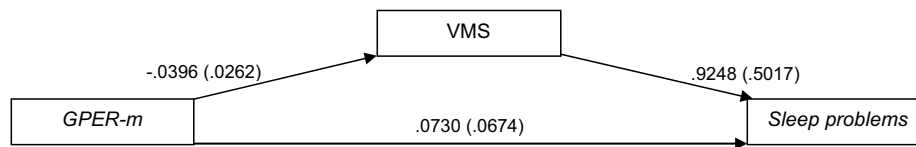


Figure 4 Path diagram of the mediation model (Macro Model 4),⁶² illustrating the associations between *GPER* methylation (*X*) and sleep disturbances (*Y*) through *VMS* (*M*). Estradiol (*E2*), menopausal status (*MS*), depressive symptoms, body mass index (*BMI*), age and caffeine consumption were included in the model and controlled in each of the three paths (a_i , b_i , c'). Values outside parentheses= unstandardized b coefficient; values in parentheses= standard error.

Abbreviations: *GPER-m*, G protein-coupled estrogen receptor methylation; *VMS*, vasomotor symptoms.

GPER methylation on the global scores of perceived sleep quality, sleep efficiency and daytime dysfunction was not significant ($b = 0.038$, $t(23) = 0.84$, 95% CI [-0.0544, 0.1295]), and nor was the indirect effect through *VMS* ($b = -0.023$, 95% CI [-0.0722, 0.0287]).

Discussion

In the present study, we examined the associations between *ESR1/GPER* methylation and sleep problems in healthy women aged 40–73 years. No direct associations between *ESR1/GPER* methylation and sleep problems were detected. However, we found that increased levels

of *ESR1* methylation were predictive of increased severity of *VMS*, which in turn predicted increased sleep problems. These associations were significant even if potential cofounders such as age, *MS*, *E2*, and medical conditions were controlled.^{3,11,44,54-56,63}

The idea that epigenetic modifications of ER genes may be key drivers of the onset and emergence of perimenopausal symptoms, such as *VMS*, has been previously suggested.²¹ One potential mechanism contributing to altered epigenetic patterns of ER genes may be the lifetime action of social and environmental determinants. For instance, early exposure to social stressors and environmental factors (eg: chemicals,

Table 4 Results of the Mediation Model with *GPER* Methylation as Predictor, 95% Bias-Corrected Confidence Interval Predicting Sleep Quality

Outcome Variable: <i>VMS</i>	<i>b</i>	<i>se</i>	<i>t</i>	<i>p</i>	LLCI	ULCI
<i>GPER-m</i> (a_i)	-.0396	0.0262	-1.5117	0.1437	-.0937	0.0145
CV						
Age	-.0749	0.0353	-2.1208	0.0445	-.1477	-.0020
<i>MS</i>	0.7074	0.3276	2.1596	0.0410	0.0313	1.3836
<i>BMI</i>	-.0210	0.0245	-.8555	0.4007	-.0716	0.0296
<i>E2</i>	-.0478	0.0414	-1.1542	0.2598	-.1332	0.0376
Depressive symptoms	-.0228	0.0304	-.7485	0.4614	-.0856	0.0400
Caffeine	0.2073	0.1481	1.3999	0.1743	-.0983	0.5129
Outcome variable: sleep problems (PSQI)						
<i>VMS</i> (b_i)	0.9248	0.5017	1.8434	0.0782	-.1131	1.9627
CV						
Age	0.1699	0.0945	1.7971	0.0855	-.0257	0.3655
<i>MS</i>	-1.2543	0.8799	-1.4256	0.1674	-3.0746	0.5659
<i>BMI</i>	-.0036	0.0612	-.0594	0.9531	-.1302	0.1229
<i>E2</i>	0.0183	0.1045	0.1751	0.8625	0.1978	0.2344
Depressive symptoms	0.2018	0.0756	2.6686	0.0137	0.0454	0.3583
Caffeine	-.6193	0.3785	-1.6360	0.1155	-1.4023	0.1638
Direct/indirect effects on sleep problems (PSQI)						
Direct effect (c')	0.0730	0.0674	1.0834	0.2899	-.0664	0.2124
Indirect effect ($a_i b_i$)	-.0366	0.0367			-.1083	0.0416

Notes: a_i represents the association between *GPER-m* and *VMS*; b_i represents the association between *VMS* and sleep quality; c' represents the direct effect of *GPER-m* on *Y*, after adjusting for all other variables included in the model; $a_i b_i$ represents the indirect effect of *GPER-m* on sleep quality through *VMS*; b represents the **unstandardized regression coefficient**.

Abbreviations: *se*, standard error; BootLLCI, bootstrapping lower limit confidence interval; BootULCI, bootstrapping upper limit confidence interval; *VMS*, vasomotor symptoms; *GPER-m*, G protein-coupled estrogen receptor gene methylation; CV, control variables; *MS*, menopausal status; *BMI*, body mass index; *E2*, estradiol.

nutrition) has been linked to altered *ESR1* methylation.^{6,23,33,64} Most interestingly, early life adversities, such as abuse and neglect, have been previously linked to increased *ESR1* methylation in rat,²³ and women,³³ as well as to increased VMS among midlife women.⁶⁵

The Study of Women Across the Nation (SWAN), assessing sleep quality in a multi-ethnic sample of 3'045 women in the US, showed that the prevalence of sleep disorders ranged from 16% to 42% in premenopausal women, from 39% to 47% in perimenopausal women, and from 35% to 60% in postmenopausal women, with Caucasian women reporting the highest rates.⁶⁶ In our study, sleep problems with PSQI scores >5 were reported by 24% of premenopausal women, 31.3% of perimenopausal women and 35.5% of postmenopausal women. The characteristics of our entirely healthy sample may explain the slightly lower prevalence of poor sleep identified in the present study.¹¹ This may also contribute to explain the smaller direct effect of *ESR1* methylation on sleep problems, compared to the indirect effect mediated by VMS. The prevalence of VMS among the participating women was comparable with the results reported by the SWAN, which indicated that up to 80% of peri- and postmenopausal women experience VMS.⁴⁹ In both the present study and previous studies, VMS were strongly associated with sleep problems.³ Together, these results may point to VMS as a primary cause of sleep problems also in healthy women.

The assessment of epigenetic modifications in peripheral tissues, such as blood, is increasingly seen as an important source of biomarkers in health and disease. In addition, as epigenetic signatures are mitotically heritable and yet reversible, their correction holds considerable therapeutic potential.^{67,68} The clinical implications of *ESR1* methylation in VMS may merit further investigation. For instance, longitudinal studies which assess the status of *ESR1* methylation before the perimenopausal phase, and which monitor the development of VMS, may provide information regarding the potential of *ESR1* methylation as a predictive marker of VMS. In the present study, we used DNA extracted from peripheral blood collected on dried blood spots (DBS). The DBS technology has been suggested as a promising tool in the context of research on biomarkers.⁶⁹ Its advantages include long-term storage of samples even at room temperature, and ease of transportation.⁷⁰

Although VMS may be primarily neurological symptoms involving brain regions such as hypothalamic areas, studying DNA methylation in the brain relies on post-mortem tissues, which has several implications in terms

of impracticality.^{71,72} Therefore, peripheral tissues such as blood, buccal cells and saliva are currently used as a proxy of the brain in methylation studies.^{73,74} Studies using peripheral tissues as a surrogate have demonstrated how specific methylation signatures are associated with disorders affecting the brain.⁷⁵ In addition, although DNA methylation is tissue-specific, it has been demonstrated that at some CpGs, DNA methylation variance is more closely linked to individual specificity than to tissue specificity.⁷⁶ Online tools have been developed which enable a comparison of inter-individual methylation variance between peripheral tissues and the brain.^{76–78} Using the online tool provided by Hannon et al (2015),⁷⁶ we found that methylation of CpGs located in the assessed DNA region of *ESR1* was strongly correlated between blood and brain in the same individuals.³⁵ Therefore, methylation variations in blood may reflect methylation variations in the brain, for the assessed CpGs of *ESR1*.

Strengths and Limitations

To the best of our knowledge, this is the first study to examine and highlight associations between *ESR1* methylation and sleep problems through VMS in healthy middle-aged and older women. In this study, we assessed subjective sleep through the PSQI,^{38,79} while objective parameters of sleep were not measured. Indeed, questions have been raised regarding the validity of laboratory sleep assessments in women experiencing VMS.⁸⁰ For instance, evidence supporting an association between objective measures of sleep and VMS is limited and contradictory, despite the association between sleep problems and VMS is well known.⁸¹ However, as subjective and objective sleep measurements may capture different aspects of sleep,^{80,82} objective sleep assessments, such as polysomnography or actigraphy, may be needed to better elucidate the nature of the association between ER genes methylation and sleep.

Our sample included only healthy Caucasian women. On the one hand, as both the race and health status may affect subjective sleep difficulties and self-perceived menopausal symptoms,^{3,53,66} the characteristics of our sample may help to limit biases linked to race and disease factors. On the other hand, our findings may not be generalizable to other racial/ethnic groups and to the general population including women with poor health. In addition, the sample of this study included three menopausal groups, and the distribution of participants among the three groups was not equal. In particular, the perimenopausal group included significantly fewer participants than the pre- and postmenopausal groups. These

considerations may limit comparison with other studies. Furthermore, the sample size of *GPER* methylation, which was strongly reduced compared to the sample size of *ESR1* methylation. Therefore, the assessment of *GPER* methylation in a larger sample may be useful in order to compare the relative contribution of *ESR1* and *GPER* methylation in the associations with sleep problems and VMS. Finally, the fact that we did not adjust for cell-type composition may also constitute a limitation of this study, as cell-type composition potentially affects levels of methylation in peripheral blood.⁸³

Conclusion

The main result of this study indicates that increased *ESR1* methylation may be associated with VMS, which in turn contribute to increased sleep problems in healthy middle-aged and older women. Therefore, this study supports the role of *ESR1* in middle-aged and older women's sleep problems through epigenetic mechanisms. Clinical implications, such as the potential of *ESR1* methylation as a biomarker of VMS, may be evaluated in future investigations.

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Disclosure

The authors report no conflicts of interest in this work.

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