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Sex Hormones and Calcitonin Gene–Related Peptide in Women With Migraine: A Cross-sectional, Matched Cohort Study

Bianca Raffaelli, MD[1, 2]; Elisabeth Storch[1]; Lucas Hendrik Overeem, MSc[1]; Maria Terhart[1]; Mira Pauline Fitzek, MD[1]; Kristin Sophie Lange, MD[1]; Uwe Reuter, MD[1, 3]

Corresponding Author: Bianca Raffaelli, bianca.raffaelli@charite.de

1. Department of Neurology, Charité – Universitätsmedizin Berlin, Berlin, Germany

2. Clinician Scientist Program, Berlin Institute of Health at Charité (BIH), Berlin, Germany

3. Universitätsmedizin Greifswald, Greifswald, Germany

Equal Author Contribution:

Contributions:
Bianca Raffaelli: Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Study concept or design; Analysis or interpretation of data
Elisabeth Storch: Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data; Analysis or interpretation of data
Lucas Hendrik Overeem: Drafting/revision of the manuscript for content, including medical writing for content; Analysis or interpretation of data
Maria Terhart: Drafting/revision of the manuscript for content, including medical writing for content; Major role in the acquisition of data
Mira Pauline Fitzek: Drafting/revision of the manuscript for content, including medical writing for content
Kristin Sophie Lange: Drafting/revision of the manuscript for content, including medical writing for content
Uwe Reuter: Drafting/revision of the manuscript for content, including medical writing for content; Study concept or design; Analysis or interpretation of data

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Abstract

Background and Objectives: Sex hormones may modulate CGRP release in the trigeminovascular system. We studied CGRP concentrations in plasma and tear fluid in female participants with episodic migraine (EM) and a regular menstrual cycle (RMC), female participants with EM and combined oral contraception (COC), and female participants with EM in the postmenopause. For control, we analyzed three corresponding groups of age-matched female participants without EM.

Methods: Participants with a RMC had two visits: during menstruation on menstrual cycle day 2 ± 2 and in the periovulatory period on day 13 ± 2. Participants with COC were examined at day 4 ± 2 of the hormone-free interval (HFI) and between days 7-14 of hormone intake (HI). Postmenopausal participants were assessed once at a random time point. Plasma and tear fluid samples were collected at each visit for determination of CGRP levels with an enzyme-linked immunosorbent assay.

Results: A total of 180 female participants (n=30 per group) completed the study. Participants with migraine and a RMC showed statistically significantly higher CGRP concentrations in plasma and tear fluid during menstruation compared to female participants without migraine [plasma: 5.95 pg/ml (IQR 4.37 – 10.44) vs. 4.61 pg/ml (IQR 2.83 – 6.92), p=0.020 (Mann-Whitney U test); tear fluid: 1.20 ng/ml (IQR 0.36 – 2.52) vs. 0.4 ng/ml (IQR 0.14 – 1.22), p=0.005 (Mann-Whitney U test)]. In contrast, female participants with COC and in the postmenopause had similar CGRP levels in the migraine and the control groups. In migraine participants with a RMC, tear fluid but not plasma CGRP concentrations during menstruation were statistically significantly higher compared to migraine participants under COC (p=0.015 vs. HFI and p=0.029 vs. HI, Mann-Whitney U test).

Discussion: Different sex hormone profiles may influence CGRP concentrations in people, with current or past capacity to menstruate, with migraine. Measurement of CGRP in tear fluid was feasible and warrants further investigation.

Introduction

The prevalence of migraine is three times higher in women than in men[1]. Fluctuations of sex hormones play a crucial role in the pathophysiology of the disease[2]. The estrogen-withdrawal-hypothesis suggests that a drop in estrogen plasma concentrations can trigger migraine attacks[3]. In line with this hypothesis, migraine frequency and pain severity are higher during the perimenstrual phase of the menstrual cycle but also in the perimenopausal period before hormonal stabilization at an older age[2, 4]. Migraine prevalence gradually declines after natural menopause[5].

Hormonal contraception leads to the suppression of physiological hormonal fluctuations with variable effects on migraine[6]. The most common hormonal contraception in Europe and North America are combined estrogen-progesterone oral compounds (combined oral contraceptives, COC)[7]. While some patients experience an improvement of migraine with COC, others experience worsening, with migraine attacks occurring most frequently during the seven-day hormone-free interval (HFI)[6].

The pathophysiological mechanisms leading from hormonal changes to the development of migraine attacks are complex. The neuropeptide Calcitonin Gene-Related Peptide (CGRP) has a key role in migraine initiation[8] and is likely to have a relevant function in the processes initiated by sex hormones changes. During a migraine attack, CGRP is released from trigeminal afferents and triggers an inflammatory response[9]. Preclinical research suggests that sex hormones fluctuations can lead to activation of the trigeminovascular system and subsequent release of CGRP, which may contribute to the high prevalence of migraine in female persons of childbearing age[10]. However, the clinical evidence in humans is inconclusive. While older investigations suggest a direct relationship between estrogen and CGRP concentrations[11, 12], newer studies imply a higher CGRP release in low estrogen phases[13, 14].

The accurate measurement of CGRP in peripheral blood is challenging due to its very short half-life time, degradation, and dilution effects after release[15]. A recent pilot study detected increased CGRP concentrations in tear fluid in participants with migraine compared to control participants without migraine[16]. This exploratory method is non-invasive and could provide a more direct measurement of the trigeminal CGRP release due to its spatial proximity to the trigeminal nerve.

Here, we studied CGRP concentrations in both plasma and tear fluid of female participants with migraine and female participants without mirgraine under different hormonal conditions. We aimed to assess the relationship between sex hormones and CGRP levels, and whether the presence of migraine affects this relationship. It was our hypothesis that a) female

persons with migraine display higher CGRP concentrations than female persons without migraine during the physiological menstrual cycle and b) that the suppression of naturally occurring sex hormones through COC or after menopause is associated with changes in the CGRP concentrations.

Methods

Study design and participants

This is a cross-sectional, matched-cohort study at the Headache Center, Department of Neurology, Charité Universitätsmedizin Berlin, Berlin, Germany.

The study cohort consisted of three groups of female participants with episodic migraine: 1) With a regular menstrual cycle (M-RMC); 2) Under contraceptive treatment with a COC (M-COC); 3) During the postmenopause (M-PM). For control, we studied three respective groups of age-matched control female participants without episodic migraine (C-RMC, C-COC, and C-PM).

Participants with migraine were recruited from our outpatient headache clinic. For the recruitment of participants without migraine, we contacted hospital and university staff via announcements in mailing lists or direct approach.

Inclusion and exclusion criteria

Episodic migraine was defined according to the International Classification of Headache Disorders 3 (ICHD-3)[17]. All female participants with migraine should have had at least three days with migraine in the four weeks prior to screening, as documented in a headache diary.

A RMC was defined as cycle duration of 28 ± 2 days in the three months before screening. In this group, the diagnosis of menstrually-related migraine[17] was required for study participation. For inclusion in the COC groups, female participants should confirm the regular use of the same contraceptive drug in a 21/7 regimen (i.e. 21 days of hormone intake [HI] followed by a 7-day hormone-free interval [HFI]), beginning at least three months prior to screening. For the postmenopausal groups, the last menstruation should have occurred at least 5 years before inclusion in the study.

Exclusion criteria were: any other diagnosed primary headache disorder except tension-type headache on less than 2 days in the month prior to screening; concurrent migraine preventive drug treatment; any gynecological or other neurological diseases; ophthalmologic conditions interfering with lacrimation; any other relevant diseases requiring regular medication; hormonal treatment with indications other than contraception; pregnancy;

lactation; post-sterilization. For participants with migraine and a RMC, the diagnosis of pure menstrual migraine[17] led to exclusion from the study.

Study procedures

Before the beginning of experimental procedures, potential participants were screened for eligibility. Eligible individuals had an initial interview to record their medical history and a physical examination. In participants with migraine, we reviewed their headache calendars of the month prior to screening.

The study protocol for female participants with a RMC consisted of two study visits. The first visit was scheduled at day 2 ± 2 of menstrual cycle (during menstruation), while the second visit took place at day 13 ± 2 of menstrual cycle (periovulatory period). These time intervals were selected because estrogen levels are at their lowest during menstruation and at their highest during ovulation.

Female participants with COC were assessed twice: at day 4 ± 2 of the HFI and between days 7-14 of HI. Postmenopausal female participants had only one visit at a variable time point.

All visits in participants with migraine were performed in the interictal period, defined as a state free of any migraine symptoms and free of acute pain medication for 12 hours before and after each visit. Participants were instructed to call and reschedule the appointment in case of migraine or acute medication intake within 12 hours before the scheduled visit. We also contacted all participants by phone the day after each visit and asked about any migraine symptoms or medication intake in the 12 hours after study visit. If this was the case, the visit was repeated at the next possible time point.

Sample preparation and analytical procedures

Each visit took place between 9 a.m. and 5 p.m. in a non-fasting condition. Blood and tear fluid samples were collected following standardized protocols[16, 18].

For CGRP measurement, blood was collected in precooled 4 ml EDTA tubes (BD Vacutainer®), that were previously prepared with 150 µl aprotinin (3-7 trypsin inhibitor unit (TIU)/ml) (Sigma Aldrich, Munich, Germany). The tubes were immediately centrifuged for 15 minutes at −6°C and 2000 rpm. Plasma was then transferred in 1.5 ml polypropylene tubes (Eppendorf, Hamburg, Germany). We collected tear fluid from the lateral canthus of one eye with a 10µl glass capillary (Brand™, Wertheim, Germany). In participants with migraine, we selected the eye on the side on which migraine occurred most frequently. If there was no side preference and in participants without migraine, the right side was chosen by default. The capillary was removed after reaching the maximal volume of 10µl or after 60 seconds at the latest. If the eye showed signs of irritation, such as redness or pruritus, the procedure

was stopped immediately. A lack of tear production after one minute led to exclusion from the study. The volume of tear fluid collected was determined (range: 1.4 to 10.0 µl) and tear fluid then transferred in a 1.5 ml tube containing 500 µl of tissue protein extractor solution (TPER; Pierce Rockford, IL). Both plasma and tear fluid samples were stored at -80°C. We measured CGRP concentrations in plasma and tear fluid with a commercial sandwich Enzyme-linked Immunosorbent Assay (ELISA) kit (CUSABIO®, Wuhan, China), following manufacturer's instructions. The detection range of this kit is 1.56–100 pg/ml, the minimal detectable dose 0.39 pg/ml. However, the company does not disclose the specific recognition site of the ELISA antibodies. The kit has high intra-assay and inter-assay precision (coefficients of variation < 8% and < 10%, respectively). Using this kit, mean CGRP concentrations in previous cohorts without migraine range from 4.2 pg/ml to 6.6 pg/ml in plasma[16, 19-21] and between 0.7 and 0.8 ng/ml in the tear fluid[16, 19].

Additionally, blood was collected in 5 ml serum tubes (BD Vacutainer®) at room temperature and sent to our partner laboratory (Labor Berlin, Charité Vivantes GmbH) for the analysis of sex hormones. The following hormones were assessed via electrochemiluminescence immuno-assay: estradiol, progesterone, testosterone, luteinizing hormone (LH), follicle-stimulating hormone (FSH).

Endpoints

The primary endpoint of the study was the difference in CGRP concentrations in plasma (pg/ml) between M-RMC and C-RMC. Secondary endpoints were the differences in CGRP plasma concentrations between M-COC and C-COC and between M-PM and C-PM.

The differences in tear fluid CGRP concentrations (ng/ml) between the migraine and the control groups were considered exploratory endpoints.

As further exploratory endpoints, we analyzed correlations between CGRP levels at both study visits in participants who were measured twice and assessed the differences in CGRP plasma and tear fluid concentrations among the three migraine and the three control groups. We also analyzed correlations between the estrogen and progesterone levels and the CGRP concentrations in tear fluid and plasma.

In addition, the total cohort of participants with migraine was compared with the cohort of participants without migraine.

Statistical analysis

Sample size calculation was performed using the software G*Power[22]. Based on a previous study on interictal CGRP plasma levels in patients with migraine compared to controls without migraine[23], we assumed a large effect size of d = 0.8 for the primary endpoint. A sample size of 30 participants per group was therefore sufficient to detect an effect of similar

magnitude with a statistical power of 0.80 at a significance level of α = 0.05 (two-tailed) using the Mann-Whitney U test. Similar statistical considerations apply for differences in tear fluid concentrations[16]. We therefore aimed at 30 participants per group with complete data sets.

We summarized demographic, anamnestic and laboratory data using descriptive statistics with median and interquartile ranges (IQR) for numerical variables, and frequencies and percentages for categorical variables. Given the non-normal data distribution, we compared outcomes between groups using the Mann-Whitney U test or the Kruskal-Wallis ANOVA, as appropriate. Correlations were tested using Spearman rank correlations.

Statistical analysis was performed with SPSS Statistics 27 (IBM Corp., Armonk, NY, USA). No adjustment for multiple comparisons was made for the exploratory outcome measures.

Standard Protocol Approvals, Registrations, and Patient Consents

The study protocol was approved by the Charité Ethical Committee (EA1/004/20). All participants gave written informed consent following study information.

Data availability

Data not provided in the article because of space limitations may be shared (anonymized) at the request of any qualified investigator for purposes of replicating procedures and results.

Results

Between August 2020 and May 2022, n=196 persons who self-identified as women participated in the study. Study protocol was completed by n=180 female participants, n=30 per group. Reasons for drop-out were: no sufficient lacrimation (n=11), occurrence of migraine in the 12 hours after study visits with no possible rescheduling (n=4) and lost to follow-up (n=1).

Demographic characteristics were similar between the migraine groups and the respective control groups. Table 1 shows the demographics across all groups and key migraine features in the three migraine groups. All female participants with migraine and a RMC reported migraine attacks within the perimenstrual period during most months[17].

Female participants with a regular menstrual cycle

M-RMC and C-RMC presented physiological hormonal levels at the two study visits with low estrogen concentrations during menstruation and high estrogen concentrations in the periovulatory period (Table 2). Progesterone levels were low at both time points since both visits occurred before the luteal progesterone increase (Table 2).

During menstruation, CGRP concentrations in both plasma and tear fluid were statistically significantly higher in interictal participants with migraine compared to female participants

without migraine [plasma: 5.95 pg/ml (IQR 4.37 – 10.44) vs. 4.61 pg/ml (IQR 2.83 – 6.92), p=0.020; tear fluid: 1.20 ng/ml (IQR 0.36 – 2.52) vs. 0.4 ng/ml (IQR 0.14 – 1.22), p=0.005] (Figure 1).

CGRP levels in the periovulatory period were numerically higher in female participants with migraine compared to participants without migraine but failed to reach statistical significance [plasma: 6.28 pg/ml (IQR 3.56 – 9.48) vs. 4.87 pg/ml (IQR 2.95 – 6.41), p=0.089; tear fluid: 0.70 ng/ml (IQR 0.18 – 2.29) vs. 0.63 ng/ml (IQR 0.14 – 1.22), p=0.225].

There was a strong intraindividual correlation between the CGRP concentrations in the menstrual and the periovulatory visits, both in plasma (rho = 0.809, p<0.001) and tear fluid (rho = 0.635, p<0.001).

Female participants with combined oral contraception

Both M-COC and C-COC showed suppressed concentrations of naturally occurring sex hormones. CGRP concentrations in plasma and tear fluid were similar between participants with migraine and controls without migraine during the HFI and during HI (Table 3). There was a strong intraindividual correlation between the CGRP concentrations at both visits (plasma: rho = 0.797, p<0.001; tear fluid: rho = 0.615, p<0.001).

Postmenopausal female participants

Both postmenopausal groups showed physiological hormonal profiles with high concentrations of LH and FSH and low concentrations of estrogen, progesterone, and testosterone. There was no statistically significant difference in CGRP concentrations in plasma and tear fluid between M-PM and C-PM (Table 4).

Comparison of CGRP levels in female participants with migraine in different hormonal states

Among all participants with migraine, CGRP plasma concentrations were similar among all groups and visits (p=0.195 among all groups). In the tear fluid, female participants with a RMC had statistically significantly higher CGRP concentrations during menstruation compared to female participants under COC (p=0.015 vs. HFI and p=0.029 vs. HI) (Figure 2).

There was no correlation between the absolute estrogen and progesterone concentrations and the CGRP concentrations in plasma and tear fluid (p>0.17 for all analyses).

Comparison of CGRP levels in female participants without migraine in different hormonal states

In plasma, CGRP concentrations of control female participants with a RMC were lower than those of female participants under COC treatment and postmenopausal female participants (menstruation vs. HI: p = 0.035; ovulation vs. HI: p = 0.030; menstruation vs. postmenopause: p = 0.015; ovulation vs. postmenopause: p = 0.013) (Figure 3). No statistically significant correlation between absolute sex hormone concentrations and CGRP concentrations could be detected (p>0.17 for all analyses). CGRP levels in the tear fluid were similar across all groups and all visits of control female participants (p=0.622 among all groups).

CGRP plasma vs. tear fluid measurements

Across all subjects (n=180) and study visits (n=300), CGRP concentrations were 5.48 pg/ml (3.98-7.82) in plasma and 0.51 ng/ml (0.16-1.22) in tear fluid. Tear fluid concentrations were 80.5x higher than in plasma (IQR 27.8 – 260.7).

Overall, participants with migraine had statistically significantly higher CGRP levels in tear fluid compared to participants without migraine [migraine groups: 0.67 ng/ml (IQR 0.17 – 1.59) vs. control groups: 0.41 ng/ml (IQR 0.15 – 0.80), p = 0.013]. Plasma concentrations were similar with 5.22 pg/ml (IQR 4.03-7.97) in the migraine groups vs. 5.95 pg/ml (IQR 3.73 – 7.79) in the control groups (p = 0.965).

Discussion

CGRP levels in plasma and tear fluid in this large cohort of female participants varied depending on the presence of migraine and the hormonal status. Female participants with episodic migraine had higher interictal CGRP concentrations in plasma and the tear fluid during menstruation than female participants without migraine. This finding did not apply to female participants with COC and during the postmenopause. In female participants with migraine, the suppression of the hormonal fluctuations through COC treatment was associated with lower CGRP tear fluid levels than during physiological menstruation.

Our findings suggest a link between sex hormones and CGRP in migraine pathophysiology in humans. The influence of sex hormones – in particular estrogen - on intracranial CGRP release has been studied mainly in vitro or animal research. Estrogen receptors are highly expressed in CGRP-positive neurons in the trigeminovascular system[24] and hormonal fluctuations can modulate their excitability[10, 25]. In animal models, deficiency of female sex hormones increases CGRP expression in various brain regions[26-28]. Also in the trigeminal

ganglion, the fall of endogenous estrogen levels in ovariectomized rats led to a significant increase in CGRP expression, which decreased following estrogen replacement treatment[29]. These observations are in line with our results in female patients with migraine: the physiological estrogen drop in the perimenstrual period was associated with higher CGRP concentrations than under hormonal contraceptive treatment.

A higher CGRP release during menstruation could help to explain the biological predisposition for more frequent, severe, and long-lasting migraine attacks in this period[30]. In line with this hypothesis, menstrual migraine attacks were more frequent and severe than non-menstrual attacks even in female persons treated with the CGRP-receptor antibody erenumab[31]. Krause et al. (2021) hypothesized that a decline in estrogen levels may lead to an increased CGRP signaling and generate a pro-migraine state with an increased susceptibility for migraine attacks[25]. Of note, this seems to apply only for a decrease in naturally occurring estrogen concentrations coming from a previously higher level but not for stable low concentrations during the postmenopause. In addition, the absolute hormone concentrations do not seem to play a relevant role, but rather the changes in hormonal levels. Accordingly, all correlation analyses between estrogen or progesterone levels and CGRP concentrations did not reveal any statistically significant result.

A few older studies showed that sex hormones might affect CGRP concentrations also in individuals without migraine. Stevenson et al. (1986) detected increased concentrations of immunoreactive CGRP in plasma during pregnancy, which decreased after delivery[11]. In a pivotal study by Valdemarson et al. (1990), CGRP plasma levels were significantly higher in eleven female participants taking an oral contraception than in twelve female participants without hormonal treatment[12]. The study did not provide data on the day of menstrual cycle or the regimen of hormonal intake[12]. In accordance with these results, in our study, oral contraception in female participants without migraine was associated with higher levels of CGRP in plasma but not in the tear fluid compared to fertile female participants without contraception. The intake of exogenous hormones seems to induce systemic changes in CGRP concentrations[10], while intracranial CGRP levels as indirectly measured in the tear fluid seem to be not affected. Indeed, high estrogen states like pregnancy have been demonstrated to increase CGRP concentrations in other anatomical regions such as the spinal cord[32]. Estrogen substitution in rats led to a CGRP increase in the mesenteric arterioles, dorsal root ganglia[33, 34], and in the gastric tract[35]. Progesterone treatment induced an increased expression of CGRP receptors in the murine uterus and mesenteric arteries[36, 37]. The postmenopause is also associated with an increase in systemic CGRP levels[38], a finding which we could reproduce in our cohort of control female participants. The cardiovascular system has been proposed as the source of the elevated CGRP

concentrations, as postmenopausal female persons with vasomotor symptoms appear particularly affected[39, 40]. Taken together, hormone dependent CGRP changes in plasma of female persons without migraine seem to originate from sources other than the trigeminovascular system.

CGRP concentrations in plasma are influenced by a multitude of factors and allow limited conclusions about the release from the trigeminal nerve system[15]. It is estimated that only one fifth of CGRP in peripheral blood derives from trigeminal sources[16]. While the crucial role of CGRP in migraine pathophysiology is indisputable, the feasibility of plasma CGRP as a biomarker of migraine remains a matter of debate[15]. Previous research reported controversial results regarding interictal plasma CGRP levels in patients with episodic migraine: While some studies detected higher CGRP levels in cubital vein blood outside of acute migraine attacks, others observed no difference to controls without migraine[23, 41-43]. Our results provide a differentiated view depending on the hormonal status of the patients. Female participants with episodic migraine during menstruation had higher interictal plasma CGRP concentrations than female participants without episodic migraine, while this was not the case in the other hormonal conditions examined.

Biomaterials closer to the trigeminal CGRP source such as tear fluid may represent a more direct and suitable approach[16]. Kamm et al. (2019) reported, in n=30 interictal mix-sexed patients with episodic migraine, higher CGRP concentrations than in n=48 controls without episodic migraine[16]. In the current analysis, we could confirm and expand these findings to a significantly larger cohort. Similar to this previous study, CGRP levels in the tear fluid were much higher than in plasma possibly due to lower proteolytic activity in this liquid than in plasma. In fact, in individuals without ophthalmologic conditions, the levels of peptidases are generally low in the tear fluid[44-46]. On the contrary, CGRP in plasma is quickly sheared into shorter fragment by endopeptidases[47], which may in part explain the lower CGRP concentrations detected with a commercial ELISA. More complex methods such as high-performance liquid chromatography (HPLC) are able to detect and differentiate between different peptide fragments[47].

CGRP in the tear fluid originates mainly from trigeminal nerve fibers in the cornea and conjunctiva, while ocular autonomic nerve fibers and the lacrimal and meibomian glands express only little or no CGRP[48, 49]. Averaged over the whole cohort, the median CGRP concentrations in the tear fluid of interictal patients with migraine were higher than in controls without migraine. This corroborates the hypothesis of an increased activation of the trigeminovascular system even outside the acute attacks. However, in the analysis by subgroups, statistical significance was confirmed only in menstruating persons. Future studies should therefore take the hormonal status of the participants into account when

examining CGRP in migraine. Despite these promising findings, CGRP determination in the tear fluid lacks validation and should be considered as an exploratory procedure. For further use, a thorough validation study needs to be performed in order to compare performance characteristics of CGPR levels in the tear fluid with the current standard measurement in plasma.

This is a comprehensive analysis about sex hormones and CGRP concentrations in female persons with migraine. The three groups of female participants with migraine were similar regarding migraine frequency and intensity. The selection of age-matched female participants without migraine and without other significant diseases or regular medication represents a key strength of this investigation. The measurement of sex hormone concentrations at each visit ensured that participants were in the predefined hormonal phase. Without a continuous hormonal measurement, however, we cannot determine whether the periovulatory visits took place exactly on the day of ovulation or rather in the few days before or after. Of note, we excluded female persons with a pure menstrual migraine, who might possibly have an even stronger influence of hormonal fluctuations on migraine-inducing mechanisms. Moreover, we included only cisgender women. Therefore, the findings do not generalize to all women (e.g. transgender women). One further limitation is the definition of the interictal state, i.e. at least 12 hours free of migraine and acute medication before and after each visit. This is shorter than in other similar investigations[16]. We rationalized that the shortening of this period reduces organizational visit changes and thereby dropouts. Twelve hours are more than two elimination half-lives of most triptans and NSAIDs and we did not expect any relevant residual efficacy after this time[50]. CGRP measurement requires strict preanalytical sample handling and CGRP concentrations may vary between studies depending on the exact methodology. In this study, we followed the protocol by Kamm et al. (2019) with the most sensitive commercial ELISA kit that is available. Indeed, we found similar concentrations of CGRP in both plasma and tear fluid as described in this previous study and other studies with the same commercial kit[16, 19-21]. The detection of a strong correlation of CGRP levels between study visits in participants that were assessed twice proves a high interindividual consistency. Importantly, multiple physiological and pathological processes can influence both CGRP and sex hormone concentrations. Despite careful selection of subjects and standardized visits, we could not control for all possible confounding factors. This study is intended as a pilot study. It provides first evidence of an association between CGRP and different sex hormone profiles in humans and sets the context for further studies with lager sample sizes and adequate power to correct for multiple testing and confounders.

Conclusion

In conclusion, our data suggests hormone dependent changes in CGRP concentrations in female patients with episodic migraine. The elevated CGRP release from the trigeminovascular system following hormonal fluctuations could help to explain a higher susceptibility for migraine in female people who menstruate. The lower CGRP tear fluid concentrations under hormonal contraception in patients with migraine could be associated with an altered migraine susceptibility under hormonal therapy and should be further investigated in a longitudinal design.

Editors' Note

Neurology recognizes that sex and gender are not interchangeable. *Neurology* editors aim to ensure that papers accurately describe and report which of these variables was evaluated in a study. In this case, the authors included only female participants, and this is the terminology used throughout the paper. We were unable to find an equivalent term to use in the title, as style guidelines suggest against using "females" as a noun. Since all the participants also identified as women, we made an editorial decision to use women in the title. *Neurology* strives to affirm persons of all genders and recognizes that the findings of this article may not pertain to all persons who identify as women.

- Rebecca Burch, MD; Roy H. Hamilton, MD, MS; Holly E. Hinson, MD, MCR

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Table 1: Description of the study population.

	M-RMC	C-RMC	M-COC	C-COC	M-PM	C-PM
Age (years)	26.50 (24.00-30.00)	26.00 (24.00-31.00)	25.00 (22.75-30.00)	27.00 (22.75-31.00)	57.50 (55.75-60.00)	58.50 (55.75-61.25)
Height (m)	1.69 (1.63-1.74)	1.70 (1.63-1.72)	1.68 (1.65-1.71)	1.69 (1.63-1.74)	1.70 (1.63-1.72)	1.63 (1.60-1.67)
Weight (kg)	63.00 (53.75-73.43)	59.00 (55.00-70.75)	62.00 (56.75-70.25)	59.00 (55.00-70.75)	70.00 (60.75-77.25)	73.50 (62.00-80.50)
Cycle length (days)	28 (27-30)	28 (26-30)				
Estradiol dose in COC (mg)			0.03 (0.03-0.03)	0.03 (0.03-0.03)		
Progesterone dose in COC (mg)			2.00 (0.15-2.00)	2.00 (0.15-2.00)		
Age at menopause (years)					50.00 (48.87-51.00)	50.00 (48.75-52.00)
Age at migraine begin (years)	16.75 (12.37-22.50)		20.00 (17.75-22.13)		20.50 (15.62-31.25)	
Aura (n, %)	11, 36.7%		17, 43.3%		9, 30.0%	
Monthly migraine days	4.00 (3.87-6.25)		5.80 (4.0-7.0)		5.25 (4.00-9.00)	
Pain intensity (0-10 NAS)	7.5 (7.0-8.0)		8.0 (6.0-9.0)		7.0 (6.0-10.0)	
Attack duration (hours)	24.00 (12.00-36.00)		27.00 (9.25-48.00)		36.25 (15.75-63.00)	
Positive family history (n, %)	22, 73.3%		18, 60.0%		22, 73.3%	

Values are median (IQR) or n, %. COC = combined oral contraception. NAS = numeric analogue scale. M = female participants with migraine. C = control female participants without migraine. RMC = regular menstrual cycle. COC = combined oral contraception. PM = Postmenopause.

Table 2: Concentrations of sex hormones in participants with migraine and control participants with a regular menstrual cycle.

	Menstrual		Periovulatory	
	M-RMC	**C-RMC**	**M-RMC**	**C-RMC**
Day of menstrual cycle	3 (2-4)	2.5 (2-3)	14 (13-15)	14 (12.75-15)
Estradiol (pmol/l)	136.50 (118.75-175.75)	135.00 (99.92-169.25)	576.50 (303.00-961.25)	607.50 (320.75-1019.75)
Progesterone (nmol/l)	0.80 (0.40-1.12)	0.85 (0.50-1.32)	0.85 (0.40-2.42)	0.95 (0.47-2.72)
Testosterone (µg/l)	0.27 (0.18-0.36)	0.24 (0.14-0.34)	0.34 (0.24-0.44)	0.35 (0.21-0.47)
LH (U/l)	5.60 (4.20-6.45)	5.55 (4.00-7.30)	12.35 (7.45-31.95)	15.40 (10.67-30.72)
FSH (U/l)	5.80 (4.72-6.92)	5.80 (4.47-7.22)	6.15 (4.27-9.00)	6.45 (4.57-9.60)

Values are median (IQR). M = female participants with migraine. C = control female participants without migraine. RMC = regular menstrual cycle. LH = luteinizing hormone. FSH = follicle-stimulating hormone.

Table 3: Concentrations of sex hormones and CGRP in participants with migraine and control participants with COC treatment.

	Hormone-free interval		Hormone intake	
	M-COC	**C-COC**	**M-COC**	**C-COC**
Day of HFI / HI	3 (2-4.25)	3 (3-4)	10 (8-12)	10 (9.75-12)
Estradiol (pmol/l)	47.65 (20.27-99.70)	21.90 (18.40-58.00)	38.00 (18.40-65.15)	21.30 (18.40-46.03)
Progesterone (nmol/l)	0.30 (0.20-0.50)	0.25 (0.20-0.62)	0.35 (0.20-0.45)	0.40 (0.20-0.70)
Testosterone (µg/l)	0.15 (0.10-0.31)	0.20 (0.13-0.28)	0.14 (0.10-0.23)	0.19 (0.12-0.28)
LH (U/l)	3.20 (0.40-5.32)	1.70 (0.30-4.20)	2.60 (1.20-4.52)	2.15 (0.30-4.90)
FSH (U/l)	3.80 (1.27-7.95)	2.80 (0.30-6.07)	2.55 (1.75-4.12)	1.75 (0.30-4.52)
CGRP in plasma (pg/ml)	4.87 (4.22-6.15)	6.67 (3.76-8.56)	4.92 (3.89-6.24)	6.03 (4.40-9.42)
	p = 0.165		p = 0.099	
CGRP in tear fluid (ng/ml)	0.46 (0.10-1.01)	0.36 (0.14-0.59)	0.32 (0.09-1.44)	0.40 (0.13-0.82)
	p = 0.574		p = 0.690	

Values are median (IQR). M = female participants with migraine. C = control female participants without migraine. COC = combined oral contraception. HFI = hormone-free interval. HI = hormone intake. LH = luteinizing hormone. FSH = follicle-stimulating hormone.

Table 4: Concentrations of sex hormones and CGRP in participants with migraine and control participants without migraine during the postmenopause.

	M-PM	**C-PM**
Estradiol (pmol/l)	22.80 (18.40-52.30)	28.30 (18.40-47.32)
Progesterone (nmol/l)	0.20 (0.20-0.32)	0.20 (0.20-0.20)
Testosterone (µg/l)	0.11 (0.10-0.19)	0.10 (0.03-0.13)
LH (U/l)	36.10 (28.65-49.77)	37.40 (30.40-44.73)
FSH (U/l)	69.05 (58.70-97.25)	75.70 (61.42-104.25)
CGRP in plasma (pg/ml)	5.24 (3.89-7.14)	6.70 (5.48-8.02) p = 0.060
CGRP in tear fluid (ng/ml)	0.70 (0.34-1.50)	0.43 (0.21-1.01) p = 0.280

Values are median (IQR). M = female participants with migraine. C = control female participants without migraine. PM = postmenopause. LH = luteinizing hormone. FSH = follicle-stimulating hormone.

Figure legends

Figure 1: **CGRP concentrations in tear fluid (A) and plasma (B) in participants with migraine and control participants with a regular menstrual cycle (RMC).** M = female participants with migraine. C = control female participants.

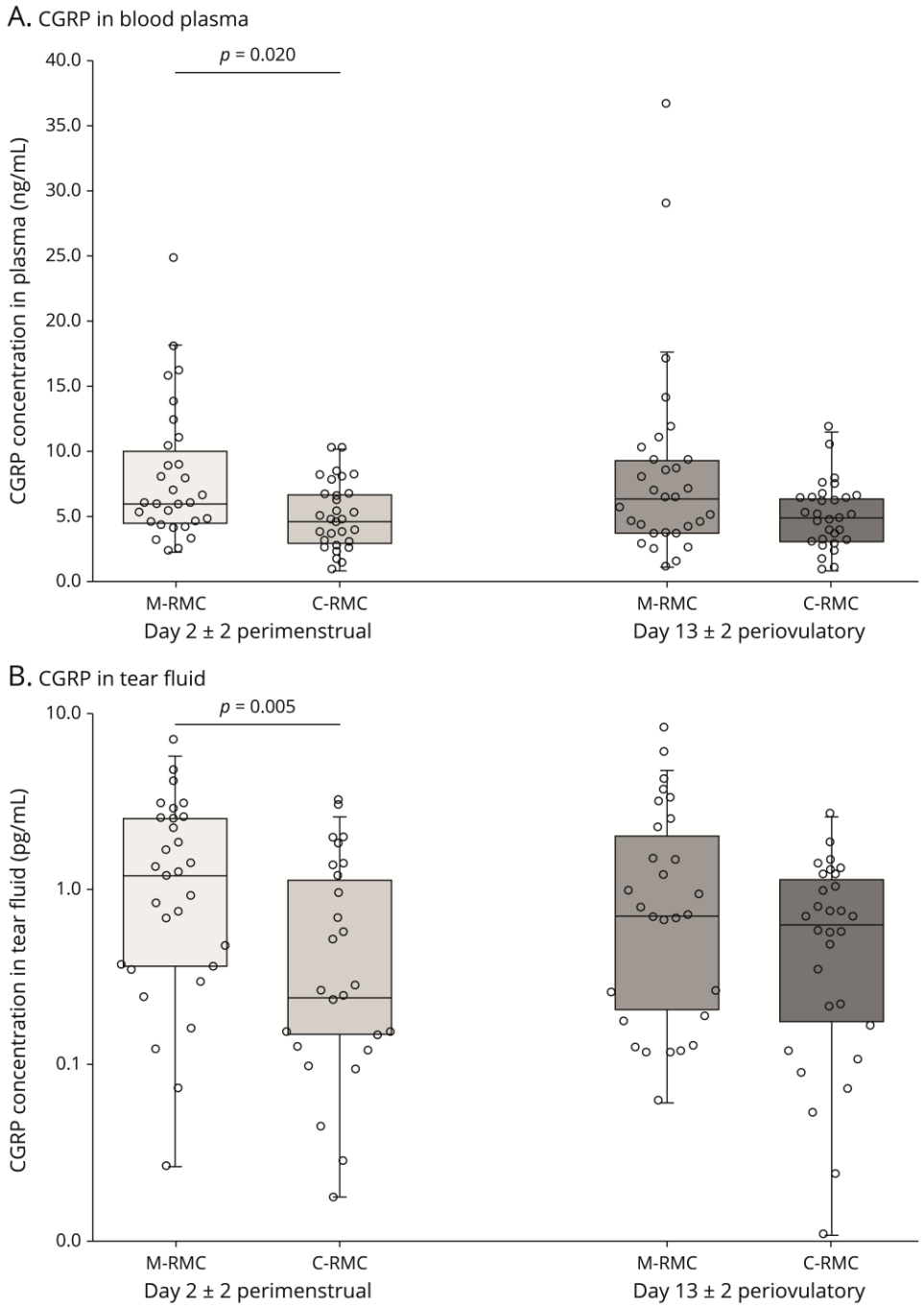

Figure 2: **CGRP tear fluid concentrations in female participants with migraine in different hormonal states.** RMC = regular menstrual cycle. COC = combined oral contraception. HFI = hormone-free interval. HI = hormone intake. PM = postmenopause.

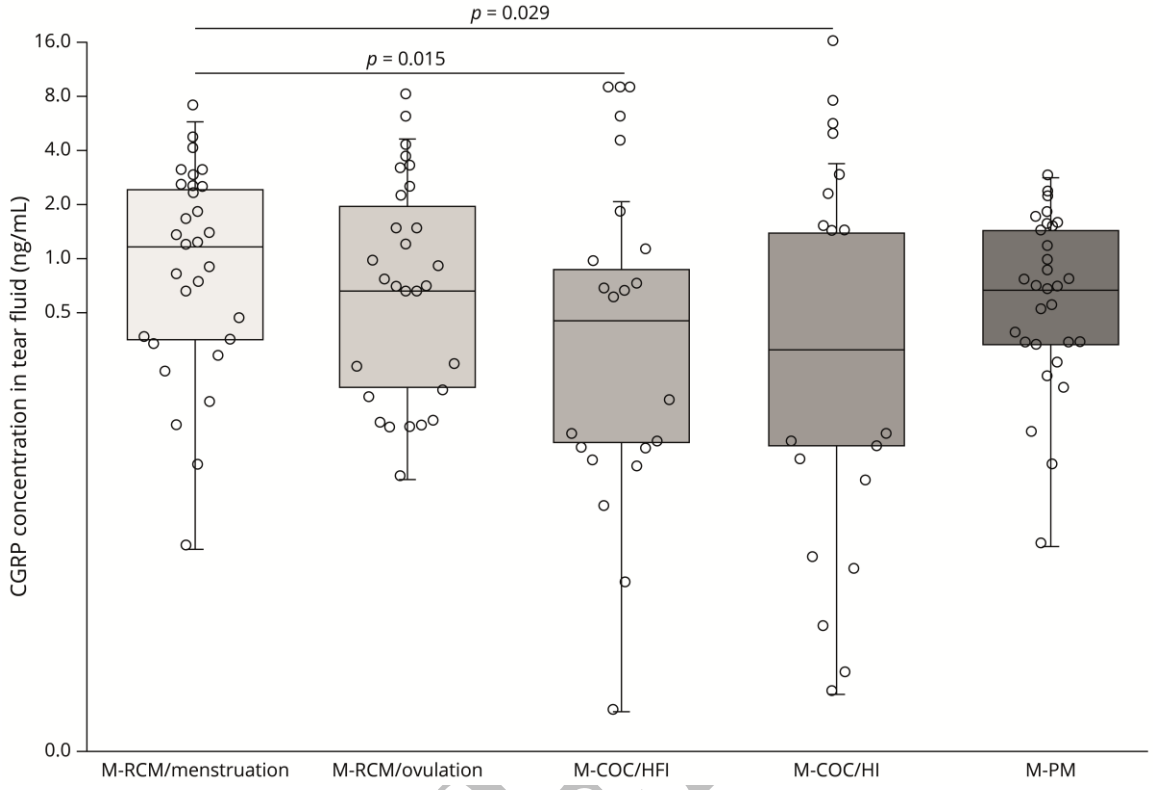

Figure 3: **CGRP plasma concentrations in female participants without migraine in different hormonal states.** RMC = regular menstrual cycle. COC = combined oral contraception. HFI = hormone-free interval. HI = hormone intake. PM = postmenopause.