Effects of *Elsholtzia splendens* and *Cirsium japonicum* on premenstrual syndrome

Mi-Sook Chung and Gun-Hee Kim

Department of Food and Nutrition, Duksum Women’s University, Ssangmun-dong, Dobong-gu, Seoul 132-714, Korea

Abstract

Premenstrual syndrome is a common chronic disorder in most women of reproductive age. The main symptoms are depression, anxiety, tension, feeling out of control, and mastalgia. In premenstrual syndrome, the effects of aromatic edible *Elsholtzia splendens* and *Cirsium japonicum* were investigated for over 3 months in 30 women participants in their twenties. In the *Elsholtzia splendens* capsule treated group, scores of depression and anxiety were significantly lower than those in the *Cirsium japonicum* capsule treated group. Moreover, instability of the premenstrual assessment form was significantly decreased in the *Elsholtzia splendens* capsule treated group. Our results suggest that *Elsholtzia splendens* could be an effective plant material in relieving symptoms of premenstrual syndrome.

Key Words: Premenstrual syndrome, depression, *Elsholtzia splendens*, *Cirsium japonicum*

Introduction

Premenstrual syndrome (PMS) is a common chronic disorder that can affect women's interpersonal relationships, social behavior, work absenteeism, and work productivity [1]. Most women in the reproductive age have one or more emotional or physical symptoms in the premenstrual phase of the menstrual cycle. The symptoms are often mild, but 5-8% has severe symptoms that are associated with substantial distress and functional impairment. Mood and behavioral symptoms, including irritability, tension, depressed mood, tearfulness, and mood swings are the most distressing, but somatic complaints such as breast tenderness and bloating can also be problematic [2-4]. In medical reports, clinically significant premenstrual symptoms were named premenstrual syndrome (PMS) or premenstrual tension (PMT). The World Health Organization's International Classification of Disease (ICD) includes “Premenstrual Tension Syndrome” under the heading of “disease of the genitourinary tract”.

For managing premenstrual syndrome, various treatments were suggested: cognitive behavioral therapy, lifestyle change, dietary supplement (Vitamin B6, calcium), herbal/complementary therapies, estradiol patches, and oral progestogens therapy [5-10]. Patients who seek medical help for PMS symptoms have usually implemented lifestyle changes and tried most over-the-counter remedies. However, before any medical treatments are commenced, it is still important to confirm that all appropriate lifestyle changes have been made. Recently, traditional herbal medicine was evaluated for relieving premenstrual syndrome [10].

*Elsholtzia splendens* is a Korean perennial, aromatic, and medicinal plant which belongs to the Labiatae family. Because *Elsholtzia splendens* contains volatile oil and flavonoids, the edible flower and leaves of this plant are primarily used for perfume and medicinal purposes [11]. Previous studies reported that *Elsholtzia splendens* had effects on reducing inflammation and fever [12].

*Cirsium japonicum*, a plant grown world-widely, is a perennial and medicinal plant which belongs to the Compositae family. The compounds most often reported for *Cirsium japonicum* are apiigenin, cirsimartin, kaempferol, linarin, luteolin, pectolinarin, and quercetin. Among them, linarin and pectolinarin are known to have anti-inflammatory properties [13]. Recently, the antidepressant effect of *Cirsium japonicum* was reported using an open field test and a forced swimming test [14]. Among the different plant parts of *Cirsium japonicum*, the leaves had the highest level of total flavonoids and apiigenin content followed by the flowers and stems [15].

Considering the aforementioned effects (reducing pain and antidepressants effects) of *Cirsium japonicum* and *Elsholtzia splendens*, it seemed appropriate to evaluate their effects on relieving premenstrual syndrome. Hence, the objective of this study was to evaluate the effects of aromatic edible *Elsholtzia splendens* and *Cirsium japonicum* on premenstrual syndrome.

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Corresponding Author: Gun-Hee Kim, Tel. 82-2-901-8496, Fax. 82-2-901-8372, Email. ghkim@doksan.ac.kr

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Subjects and Methods

Formula for capsules of Elsholtzia splendens and Cirsium japonicum

Samples of the aerial parts of _Elsholtzia splendens_ and _Cirsium japonicum_ were collected from Gyeonggi-do in the Republic of Korea. The plants were identified and authenticated by a botanist at the Korea National Arboretum. Samples were freeze-dried (FD5505, IIShin LabTech, Seoul, Korea) and the leafless woody stalks were discarded. The dried samples were subsequently crushed for 10 sec using a blender (HMC-400T; Hanil Electronics, Seoul, Korea), and then capsules were made as shown in Table 1. According to our previous experiment of total flavonoids and apigenin, the acceptable daily intake of both samples was 120 mg.

Subjects

To recruit female participants for this study, in November 2006 we advertised to students at Duksum Women's University in Seoul, Korea, entitling the study, “PMS Management Program”. All applicants were obliged to participate in an interview with this being the psychological assessment tool for screening. Thereafter, a total of 30 participants with PMS were selected according to 1) age 2) had missed menstruation cycles for more than 3 terms 3) had taken the contraceptive pill, or 3) were evaluated as unsuitable participants for this program because of serious mental illness as per SCL-90-R (Symptom Checklist-90-Revision).

Thirty participants were assigned to 3 groups, _Elsholtzia splendens_ group (n = 10), _Cirsium japonicum_ group (n = 10), and placebo group (n = 10) according to 1) age 2) Premenstrual Assessment Form-Past Cycle Version (PAF-PCV) were utilized to assess changes of emotional, behavioral and physical conditions for premenstrual duration and distinguishing between clinically anxious people and the anxiety of psychiatric patients as well. Score ranges from 20 to 80, with the higher score indicating more serious anxiety. The Cronbach's α (reinstability coefficient) of the STAI in this study was 0.61.

Table 1. Formula for capsules of _Cirsium japonicum_ and _Elsholtzia splendens_

| Material                                      | Contents (mg) | Percentage (%) |
|-----------------------------------------------|---------------|----------------|
| Freeze-dried powder of _Cirsium japonicum_ or _Elsholtzia splendens_ | 120.00        | 60.00          |
| Lactose                                       | 73.60         | 36.50          |
| Gelatin                                       | 3.52          | 1.76           |
| Carboxymethyl cellulose-Na                    | 0.88          | 0.44           |
| Magnesium stearate                            | 2.00          | 1.00           |
| Total                                         | 200.00        | 100.00         |

Table 2. Demographics characteristics

| Variables                | _Elsholtzia splendens_ group | _Cirsium japonicum_ group | Placebo group | Total |
|--------------------------|-------------------------------|---------------------------|---------------|-------|
| Sex (female)             | 10                            | 10                        | 10            | 30    |
| Age (yr)                 | 23.00 ± 0.94                  | 23.10 ± 1.79              | 22.60 ± 1.65  | 22.90 ± 1.47 |
| Menarchial age           | 13.70 ± 1.83                  | 13.10 ± 1.66              | 13.40 ± 0.70  | 13.40 ± 1.45 |
| Menstrual cycle          | 29.90 ± 2.23                  | 30.50 ± 2.88              | 32.20 ± 4.21  | 30.87 ± 3.26 |
| Menstrual duration       | 5.80 ± 1.03                   | 6.00 ± 1.05               | 6.90 ± 1.37   | 6.23 ± 1.22  |

Values are the mean ± SD.

In this study, the PAF and Unipolar Summary Scales of the PAF-PCV were utilized to assess changes of emotional, behavioral and physical conditions for premenstrual duration

Table 3. The subordinate symptoms of the premenstrual assessment form (PAF)

| The subordinate factors of PAF |
|-------------------------------|-----------------------------|
| 1. Low mood / Loss of pleasures | 10. Impulsiveness          |
| 2. Endogenous depressive features | 11. Organic mental discomfort |
| 3. Instability                 | 12. General physical discomfort |
| 4. Atypical depressive features | 13. Sign of water retention |
| 5. Hysteroid features          | 14. Autonomic physical changes |
| 6. Hostility / anger           | 15. Fatigue                 |
| 7. Social withdrawal           | 16. Impaired social functioning |
| 8. Anxiety                     | 17. Miscellaneous mood / behavior |
| 9. Increased well-being        | 18. Miscellaneous physical changes |
[6,19]. The PAF is a measure for retrospectively evaluating the PMS symptoms of menstrual duration during the previous 3 cycles and the PAF-PCV is a retrospective measure for evaluating PMS symptoms of menstrual duration in the past 1 cycle after the treatment. This scale is a 95-item self-report measure and the total score ranges from 95 to 570 with the highest score indicating more serious symptoms. The PAF emphasized the variations of symptoms from 14–17 days before menstruation to the next period of menstruation. This scale in our study consisted of 18 subordinate symptoms (Table 3). The Cronbach’s a was 0.94.

Statistical analysis

All data are expressed as mean ± standard deviation values. Statistical significance was evaluated by ANOVA and the Scheffe test (SPSS 14.0 version). Differences between and among means were considered significant when the P value was 0.05 or 0.01.

Results

The comparison of pre-post scores on depression and anxiety among groups

Effect of *Elsholtzia splendens* and *Cirsium japonicum* on premenstrual syndrome was studied via three experimental groups of thirty women with an average age of 22.9 years: 1) the *Elsholtzia splendens* group 2) the *Cirsium japonicum* group and 3) the placebo group. Participants of the experiment were, in mean menarchial age, 13.40 ± 1.45 years old. In addition, their menstrual cycle was 30.87 ± 3.26 days and the menstrual duration in mean menarchial age, 13.40 ± 1.45 years old. In addition, their menstrual cycle was 30.87 ± 3.26 days and the menstrual duration was 6.23 ± 1.22 days (Table 2). According to our previous experiment of total flavonoids and apigenin, the acceptable daily intake of *Elsholtzia splendens* and *Cirsium japonicum* sample was 120 mg. The *Elsholtzia splendens* group and the *Cirsium japonicum* group were instructed to take 1 capsule per day (which contained 120 mg of *Elsholtzia splendens* or *Cirsium japonicum* respectively), every morning 30 minutes after waking up. As for the placebo group, participants took capsules which were exactly the same, except for the absence of the plant material. This experiment was conducted for 3 consecutive months. Before a statistical demonstration, this study conducted the test about the equality of variances. As a result, the dispersion of the 3 groups was equal. In addition, ANOVA was analyzed to examine if there were significant differences of the pre-post scores on depression and anxiety among the 3 groups. The results showed that three groups significantly differed from the pre-post score of depression and anxiety. The pre-post scores of depression (BDI) were decreased from 33.50 to 23.60 in the *Elsholtzia splendens* group, from 33.60 to 30.40 in the *Cirsium japonicum* group, and from 33.70 to 33.00 in the placebo group. Finally, we discovered a greater decrease in the pre-post scores in the *Elsholtzia splendens* group than in other groups (P < 0.01). The pre-post scores of anxiety were decreased from 59.40 to 48.10 in the *Elsholtzia splendens* group, but there were increases from 50.90 to 52.00 in the *Cirsium japonicum* group and from 55.60 to 57.10 in the placebo group. Finally, there were greater decreases of anxiety scores in the *Elsholtzia splendens* group than in other two groups (P < 0.05). Moreover, the result of the Scheffe test showed that the post scores of depression and anxiety in the *Elsholtzia splendens* group were decreased more than in other groups (Table 4).

The comparison of pre-post scores on total and subordinate symptoms of the Premenstrual Assessment Form (PAF)

The significant differences between the three groups on the pre-post scores of the PAF were presented in Table 5. Although statistically significant differences of the total PAF scores were not found, the total scores of the PAF were decreased from 270.20 to 176.70 in the *Elsholtzia splendens* group, from 257.30 to 185.60 in the *Cirsium japonicum* group, and from 245.20 to 207.00 in the placebo group. As a result, the total scores of the PAF in the *Elsholtzia splendens* group were not statistically significant, but presented greater decreases compare to other groups. However, the instability in subordinate factors showed a significant difference (P < 0.05). The pre-post scores of the instability were decreased from 11.50 to 6.40 in the *Elsholtzia splendens* group and from 11.80 to 8.30 in the *Elsholtzia splendens* group, and from 245.20 to 207.00 in the placebo group. Last but not least, the instability scores in the subordinate scales of the PAF in the *Elsholtzia splendens* group showed statistically more significant differences than other groups. In addition, although significant differences were not shown statistically in all scales (except for the instability), it was found out that there were changes of the mean score between the pre-post in all scales. In particular, the *Elsholtzia splendens*
Table 5. The comparisons of the total scores and subordinate scales of the premenstrual assessment form (PAF)

| Variables                        | Treatment                                      | Elsholtzia splendens group (n = 10) | Cirsium japonicum group (n = 10) | Placebo group (n = 10) | P     |
|----------------------------------|------------------------------------------------|-------------------------------------|----------------------------------|------------------------|-------|
| Total PAF scores                 | pre                                            | 270.20 ± 82.61                      | 257.30 ± 74.81                   | 245.20 ± 73.48         | 0.530 |
|                                  | post                                           | 176.70 ± 61.33                      | 185.60 ± 53.65                   | 207.00 ± 67.47         |       |
| Low mood / Loss of pleasures     | pre                                            | 29.60 ± 12.09                       | 28.30 ± 9.32                     | 29.00 ± 12.61          | 0.319 |
|                                  | post                                           | 20.20 ± 10.34                       | 18.50 ± 6.02                     | 24.40 ± 9.44           |       |
| Endogenous depressive features   | pre                                            | 15.60 ± 4.17                        | 13.70 ± 5.01                     | 14.30 ± 4.81           | 0.643 |
|                                  | post                                           | 8.90 ± 4.12                         | 9.90 ± 3.35                      | 10.50 ± 3.92           |       |
| Instability                      | pre                                            | 11.50 ± 3.81                        | 11.80 ± 2.82                     | 10.20 ± 4.02           | 0.022*|
|                                  | post                                           | 6.40 ± 1.65                         | 8.30 ± 3.06                      | 10.80 ± 4.57           |       |
| Atypical depressive features     | pre                                            | 21.30 ± 6.75                        | 21.70 ± 6.65                     | 20.40 ± 5.84           | 0.623 |
|                                  | post                                           | 14.50 ± 6.01                        | 15.60 ± 6.00                     | 17.20 ± 6.55           |       |
| Hysteroid features               | pre                                            | 16.00 ± 6.06                        | 15.90 ± 4.63                     | 14.20 ± 4.66           | 0.101 |
|                                  | post                                           | 10.30 ± 3.20                        | 10.10 ± 4.18                     | 13.80 ± 4.94           |       |
| Hostility / Anger                | pre                                            | 19.00 ± 8.03                        | 15.70 ± 5.40                     | 14.80 ± 6.30           | 0.118 |
|                                  | post                                           | 9.40 ± 3.03                         | 9.90 ± 3.64                      | 13.80 ± 7.27           |       |
| Social Withdrawal                | pre                                            | 14.90 ± 4.73                        | 14.10 ± 4.63                     | 12.40 ± 5.95           | 0.446 |
|                                  | post                                           | 7.50 ± 3.69                         | 8.20 ± 3.91                      | 9.90 ± 5.11            |       |
| Anxiety                          | pre                                            | 14.00 ± 5.62                        | 12.10 ± 3.84                     | 12.40 ± 4.03           | 0.824 |
|                                  | post                                           | 8.70 ± 4.00                         | 9.00 ± 2.87                      | 9.70 ± 4.03            |       |
| Increased well-being             | pre                                            | 5.40 ± 2.80                         | 5.80 ± 2.00                      | 5.60 ± 2.12            | 0.527 |
|                                  | post                                           | 5.50 ± 1.84                         | 5.90 ± 2.13                      | 4.90 ± 1.91            |       |
| Impulsiveness                    | pre                                            | 11.60 ± 5.80                        | 9.50 ± 3.90                      | 9.70 ± 3.95            | 0.199 |
|                                  | post                                           | 6.30 ± 2.41                         | 7.80 ± 3.12                      | 9.20 ± 4.61            |       |
| Organic mental discomfort        | pre                                            | 17.30 ± 8.74                        | 15.40 ± 4.58                     | 16.30 ± 6.77           | 0.962 |
|                                  | post                                           | 12.70 ± 7.33                        | 12.50 ± 3.57                     | 13.20 ± 5.73           |       |
| General physical discomfort      | pre                                            | 17.00 ± 5.01                        | 17.50 ± 5.68                     | 16.30 ± 6.00           | 0.208 |
|                                  | post                                           | 11.50 ± 3.06                        | 13.10 ± 5.55                     | 15.70 ± 6.38           |       |
| Sign of water retention          | pre                                            | 9.20 ± 3.12                         | 9.00 ± 4.14                      | 10.40 ± 4.65           | 0.323 |
|                                  | post                                           | 6.30 ± 3.20                         | 6.40 ± 2.55                      | 8.20 ± 3.52            |       |
| Autonomic physical changes       | pre                                            | 17.30 ± 7.12                        | 18.20 ± 7.69                     | 17.10 ± 7.20           | 0.491 |
|                                  | post                                           | 11.70 ± 4.17                        | 15.10 ± 7.48                     | 13.30 ± 6.77           |       |
| Fatigue                          | pre                                            | 14.30 ± 4.24                        | 14.30 ± 3.90                     | 14.70 ± 5.12           | 0.759 |
|                                  | post                                           | 9.50 ± 4.04                         | 9.30 ± 3.27                      | 10.50 ± 4.20           |       |
| Impaired social functioning      | pre                                            | 31.40 ± 8.98                        | 28.10 ± 8.41                     | 28.80 ± 11.94          | 0.824 |
|                                  | post                                           | 19.50 ± 7.43                        | 21.40 ± 8.97                     | 21.50 ± 7.72           |       |
| Miscellaneous mood / Behavior    | pre                                            | 30.40 ± 8.72                        | 29.00 ± 8.74                     | 27.60 ± 9.14           | 0.385 |
|                                  | post                                           | 20.30 ± 6.09                        | 19.50 ± 8.02                     | 23.90 ± 8.08           |       |
| Miscellaneous physical changes   | pre                                            | 12.80 ± 4.32                        | 16.40 ± 6.59                     | 14.80 ± 5.96           | 0.302 |
|                                  | post                                           | 9.30 ± 4.03                         | 10.80 ± 3.62                     | 10.90 ± 4.71           |       |

Values are the mean ± SD, * P<0.05, ** P<0.01

Discussion

Premenstrual syndrome is the consequence of fluctuations in the ovarian sex steroids leading to physical and behavioral symptoms. Many symptoms of premenstrual syndrome have been described: irritability, aggregation, depression, tension, anxiety, feeling out of control, bloatedness, and mastalgia are well-known symptoms [2-4,19]. In this study, depression and anxiety of the Elsholtzia splendens group were significantly decreased at the P<0.01 and P<0.05 respectively. As shown in Table 5, instability demonstrated significant decreases in the Elsholtzia splendens group after the intake of the capsules (P<0.05). Although there were no statistical significances, the total PAF scores indicated a decrease in the Elsholtzia splendens group after the intake. Our results revealed that Elsholtzia splendens is effective for decreasing depression, anxiety, and instability which are common symptoms of premenstrual syndrome. Therefore, Elsholtzia splendens is thought to be a useful plant material for relieving premenstrual syndrome.

Recent investigation into the relief of premenstrual syndrome has employed hormone and antidepressant drugs, whereas treatment using plant materials was not studied extensively. One...
randomized placebo controlled study has shown that *Agnus castus* is an effective treatment for women with premenstrual syndrome. The effects were confirmed by the women’s self-assessment and the investigator’s evaluation [7]. The extensive treatment for St. John’s wort as an antidepressant in women with severe premenstrual syndrome resulted in a significant relief of symptoms [7]. It was reported that evening primrose oil is of little value in treating premenstrual syndrome, except for the management of cyclical premenstrual mastalgia for which it is effective [2]. By utilizing *Kampo* which is Japan’s traditional herbal medicine used to treat premenstrual syndrome, women who participated in the experiment reported improvements in their condition. In these studies, however, it was not revealed which compound of the herbal medicine was dedicated to the relief of premenstrual syndrome symptoms. As for the effect of carbohydrates on premenstrual syndrome symptoms, relatively, there are detailed explanations. In a study about whether mild to moderate premenstrual syndrome could be improved with a commercial, carbohydrate-rich beverage, the profile of mood states (tension & anxiety factors) showed a significant effect (*P* < 0.05). The carbohydrate beverage probably acts by increasing the availability of tryptophan. This, in turn, may increase serotonin synthesis which reduces the mood and appetite disturbances accompanying PMS [6].

As mentioned above, our results strongly suggest that *Elsholtzia splendens* is effective for decreasing depression, anxiety, and instability which are common symptoms of premenstrual syndrome. According to our previous experiments, the amount of total flavonoid of flower and leaf of *Elsholtzia splendens* was 2.52 mg/g and 1.99 mg/g, respectively. And the content of apigenin of flower and leaf was 18 ug/ml and 1.35 ug/ml, respectively [20]. In *Cirsium japonicum*, the amount of total flavonoid of flower and leaf was 1.54 mg/g and 2.58 mg/g, respectively. In addition, the content of apigenin of flower and leaf was 0.33 ug/ml and 2.73 ug/ml, respectively [15]. From the above data, the amount of total flavonoids for *Elsholtzia splendens* and *Cirsium japonicum* is similar, but the content of apigenin of *Elsholtzia splendens* is comparatively higher than those of *Cirsium japonicum*. We expect that, based on our previous research, biologically active compounds which had these effects are presumed to be flavonoid and apigenin from *Elsholtzia splendens*. Therefore, further study should clarify if flavonoid and apigenin extracted from *Elsholtzia splendens* decrease depression, anxiety, and instability. In addition, further experiments should be done to reveal its action mechanisms.

In conclusion, effects of *Elsholtzia splendens* and *Cirsium japonicum* on premenstrual syndrome were studied through three experimental groups: 1) the *Elsholtzia splendens* group 2) the *Cirsium japonicum* group and 3) the placebo group. *Elsholtzia splendens* is significantly effective for decreasing depression, anxiety, and instability which are common symptoms of premenstrual syndrome. Therefore, *Elsholtzia splendens* may be a useful plant material for relieving premenstrual syndrome.

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