Evaluation of Formulated Herbal Syrup (Containing Fennel, Anise, and Celery) on the Letrozole-Induced Polycystic Ovary Syndrome Model

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Received 2021 November 01; Revised 2022 February 06; Accepted 2022 February 08.

Abstract

Background: Polycystic ovarian syndrome (PCOS) is a complex endocrine disorder associated with irregular menstrual cycles, hyperandrogenism, obesity, and reduced fertility.

Objectives: The present study aimed to formulate herbal syrup based on Iranian traditional medicine (ITM) and evaluate its effect on the letrozole-induced PCOS model in female rats.

Methods: The herbal syrup contains anise, fennel, and celery seed extracts. Five different formulations were made with different percentages of additive components. Quality control and stability tests were performed on the selected formulation. During the in vivo step, 6 groups of rats were evaluated: The control group (received carboxymethyl cellulose 1% as a vehicle) and the other 5 groups (received letrozole 1 mg/kg orally for 21 days). During 21 days, daily vaginal smears were examined to detect irregularities of the estrous cycle. After induction of PCOS, rats were orally administered with herbal syrup (1, 2, 4 mL/kg) or metformin (200 mg/kg) for 28 days. Moreover, body and ovarian weights, serum follicle-stimulating hormone (FSH), luteinizing hormone (LH), estradiol, progesterone, and testosterone were measured. Finally, ovarian tissues were isolated for histological examination.

Results: The best formulation of the syrup contained the plant extract (totally 10%), sugar (50%), sodium benzoate (0.1%), and potassium sorbate (0.1%). Body weight was significantly increased in all groups compared to the control group, and after treatment, a significant weight reduction was seen in the metformin and 1-mL/kg dose group. Following PCOS induction, ovary weight was significantly increased, while after treatment, it showed a significant decrease. After 21 days of letrozole administration, induction of PCOS was confirmed by the irregularities in estrous cycles and an increase in testosterone and LH levels. After treatments with the syrup, LH levels significantly decreased in all groups (P < 0.05), and serum testosterone and FSH levels significantly decreased in the 2-mL/kg dose group compared to the PCOS group (P < 0.05). Estradiol and progesterone levels significantly increased in the treatment groups in a dose-dependent manner. Histological studies of metformin and herbal syrup groups exhibited normal follicular development with fewer and smaller cystic follicles compared to the PCOS group.

Conclusions: The herbal syrup made from anise, fennel, and celery seed extracts improved serum levels of sex hormones, recovered the ovarian morphology in PCOS-induced rats, and can be a good candidate for further clinical trials.

Keywords: Polycystic Ovarian Syndrome, Iranian Traditional Medicine, Letrozole, Foeniculum vulgare, Pimpinella anisum, Apium graveolens
cardiovascular disease, and endometrial cancer (3-5).

PCOS treatment includes various aspects of the disease, such as reproductive, metabolic, and psychological features. For instance, lifestyle modifications (diets and exercise) to treat metabolic disorders (6) and pharmacologic therapies, including metformin, oral contraceptives (OCPs), and anti-androgen drugs (such as spironolactone and cyproterone acetate) (7), are used to control the disease. However, because of the adverse effects of these drugs, such as lactic acidosis (8), weight gain, cardiovascular disease, thromboembolism, and the relative low responses to the current therapies (9, 10), it is crucial to consider alternative treatments with fewer side effects and more effectiveness (11).

Today, due to the remarkable effects of phytopharmaceuticals in managing diseases, the use of these compounds is increasing (12). In Iranian traditional medicine (ITM), many special herbal medicines have been used to prevent and cure PCOS (11, 13). Thus far, many scientific studies have proved the effect of Iranian herbal drugs in PCOS, such as Apium graveolens L. (14, 15), Pimpinella anisum L. (15, 16), Vitex agnus-castus L. (17), Trigonella foenum-graecum L. (18), Cinnamomum verum J. Presl (19), and Foeniculum vulgare Mill. (20, 21).

Fennel (F. vulgare Mill.), anise (P. anisum L.), and celery (A. graveolens L.) are aromatic medicinal plants belonging to the Apiaceae family. Because of its estrogenic characteristics (chiefly attributed to the main compound anethole), fennel has been used as a galactagogue and emmenagogue, aphrodisiac, uterine tonic compound for thousands of years (22-24). Recent studies have indicated a broad spectrum of pharmacological activities of anise, such as antibacterial (25), anti-diabetic, and hypolipidemic (26) effects. Celery has been shown to effectively prevent cardiovascular diseases, lower blood pressure, and strengthen the heart (27).

2. Objectives

Based on ITM and according to recent studies conducted on the estrogenic, antiandrogenic, emmenagogue, and antioxidant effects of fennel, anise, and celery (28), the present study was designed to prepare a suitable formulation of herbal syrup containing these 3 herbs introduced by ITM manuscripts (28), as well as to evaluate the effect of this syrup on letrozole-induced PCOS in female rats.

3. Methods

3.1. Plant Material

Dried fruits of fennel, anise, and celery were purchased from a local market and identified at the Herbarium of Traditional Medicine and Materia Medica Research Center (TMRC), Shahid Beheshti University of Medical Sciences, Tehran, Iran. Herbal Market Samples (HMS) of the fruits of A. graveolens L. (No. 548), P. anisum L. (No. 549), and F. vulgare Mill. (No. 550) have been deposited at the Herbarium of TMRC.

3.2. Chemical, Reagent, and Kits

Letrozole was obtained from Iran Hormone Pharmaceutical Co, Tehran, Iran, and Metformin from Arya Pharmaceutical Co, Tehran, Iran. The other chemicals used in this study consist of Folin-Ciocalteu reagent (Merck, Germany), sodium carbonate (Merck, Germany), gallic acid (Sigma-Aldrich, Germany), and hematoxylin and eosin staining (H&E; Parastaneh, Iran). Serum 17β-estradiol and progesterone levels were measured using ELISA kit KGE014, R&D Systems (Bio-Techne, Minneapolis, MN, USA). Serum progesterone levels were measured using Mouse/Rat Progesterone ELISA kit (Catalog Number SEI20087, Sigma-Aldrich, Germany). The serum LH level was evaluated using an ELISA kit (catalog MBS 729873 from MyBioSource, USA) and FSH level using an ELISA kit (catalog MBS 2502190 from MyBioSource, USA).

3.3. Ethical Considerations

The Ethics Committee of Shahid Beheshti University of Medical Sciences approved the proposal of the research (code: IR.SBMU.RETECH.REC.1399.899, 2021). The in vivo part of the study was done according to the NIH Animal Care and Use Committee Guide for the Care and Use of Laboratory Animals (29).

3.4. Formulation Preparation

According to the selected traditional resource (28), our chosen prescription is herbal syrup containing an aqueous extract of anise, fennel, celery seeds, and sugar. Based on the effective and maximum permitted dose of each plant, we made 5 experimental formulations containing 10% plant extract. To prepare the formulations, first, the seeds were crushed, coarsely powdered, and extracted by decoction method with distilled water (plant: water ratio 1: 10 w/v) for 30 minutes. The filtrate was evaporated and concentrated to a final volume (concentration ratio 66:10). In order to achieve an appropriate viscosity and taste of the formulation, carboxymethylcellulose (CMC) and sugar were added, and various formulations with different ratios of mentioned ingredients (F1 - F5) were prepared. Moreover, sodium benzoate and potassium sorbate were used in the syrup as antimicrobial preservatives.
3.5. Physicochemical Quality Control of the Herbal Syrup

Various physicochemical parameters, including macroscopic and organoleptic characteristics, dried residue, cap locking, density, viscosity, pH, crystallization evaluation, and microbial content, were tested on the final formulation. In addition, the total phenolics content of the herbal syrup was determined by the Folin-Ciocalteau method (30). All tests were performed on 3 batches of the final formulation.

3.5.1. Accelerated Stability Test

According to the International Conference on Harmonization (ICH) Guidelines, the stability study for the final product was completed at accelerated stability conditions (31). Three bottles of syrup were placed at 40 ± 2.0°C in an oven for 6 months. Later, samples were examined every 3 months for the above-mentioned measurements.

3.6. Experimental Protocol

3.6.1. Animals

In this study, 42 healthy female Sprague-Dawley rats aged 6-8 weeks with an average weight of 130 g were prepared from TMRC. Animals were kept in temperature-controlled rooms (22°C), with 45% - 65% humidity and 12-hour light and dark cycles, in a pathogen-free environment. The rats were allowed water and food (standard diet) ad libitum. According to the previous studies (32, 33) and conversion of the human dose to animal dose (34), the treatment groups received doses of 1, 2, 4 mL/kg.

3.6.2. Letrozole Induced-Polycystic Ovarian Syndrome

PCOS was induced by Ndeingang et al.’s method (35). By checking the vaginal smears, the estrous cyclicity of the animals was monitored, and only rats with 2 - 3 regular estrous cycles during the 12 to 14 days of vaginal smear get into the scheme. Group I served as the control group and received 1 mL of 1% CMC (vehicle). Other groups were administered with 1 mg/kg letrozole, dissolved in 1% CMC (2 mL/kg) once daily for 21 days. Vaginal smears were obtained daily to verify the induction of PCOS. Animals were randomly sorted into 6 groups (n = 7 animals in each group). One rat from each group was sacrificed after 21 days of receiving letrozole. Biochemical and histological tests were conducted to confirm PCOS in the rats. From day 22, the animals of group II were gavaged with 30% sugar in distilled water (PCOS group), the animals of group III - V were gavaged with herbal syrup at doses of 1, 2, and 4 mL/kg and served as dose 1, 2, and 3 groups, whereas the animals of group VI were treated with metformin at a dose of 200 mg/kg (36) and considered as the metformin group. Herbal syrup and metformin were given daily for 28 days. At the end of the treatment period, rats were weighed and anesthetized with ketamine/xylazine (5/1 mg/kg) (37). The blood samples were taken for hormonal analysis, and ovaries were collected for histological evaluation.

3.6.3. Serum Hormone Analysis

Blood samples were directly collected from the hearts of the subjects at the end of the experiment. Serum samples were separated by centrifugation and stored at -70°C until use. Hormone levels (FSH, LH, estrogen, testosterone, and progesterone) were measured by enzyme-linked immunosorbent assay (ELISA).

3.6.4. Histological Analysis

On day 49, both ovaries were collected from each animal, quickly removed, cleaned up, and weighed. Later, the ovary was fixed in 10% neutral-buffered formalin for 48 hours. Then, the tissues were embedded in paraffin and cut into 5 µm sections, stained by the H&E staining method. Histological assessment was conducted under light microscopy (Ceti Microscopes, UK). Follicles were identified and scored in different stages: primary, secondary, antral, Graafian, cystic follicles, and corpora lutea.

3.7. Statistical Analysis

The results of the different groups were analyzed by 1-way analysis of variance with the Tukey-Kramer multiple pair comparison test. Statistical difference was considered significant if \( P < 0.05 \). Data were displayed as mean ± SEM.

4. Results

4.1. Formulation Preparation and Physicochemical Quality Control

Five experimental formulations were prepared (Table 1). F5 had an acceptable taste and viscosity compared to other formulations and was taken as the final preparation. The results of quality evaluation tests are presented in Table 2. The quality of the syrup was acceptable based on the physical parameters. The microbial count of F5 was within the allowable limits over 6 months, according to the British Pharmacopoeia (38).

4.2. Effects of Herbal Syrup on the Weight of Body and Ovary

The body weight measurement of the rats had a significant increase before and after treatment in all groups compared to the control group. In contrast, the weight of metformin and dose 1 groups was significantly reduced compared to the PCOS group (Figure 1). Ovaries’ weight significantly increased in the PCOS group compared to the control group and decreased in groups treated with the
Table 1. Ingredients of the Experimental Formulations

<table>
<thead>
<tr>
<th>Ingredients (%)</th>
<th>F1</th>
<th>F2</th>
<th>F3</th>
<th>F4</th>
<th>F5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plant extract</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Sugar</td>
<td>30</td>
<td>30</td>
<td>30</td>
<td>40</td>
<td>50</td>
</tr>
<tr>
<td>CMC</td>
<td>0</td>
<td>0.1</td>
<td>0.2</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Sodium benzoate</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
</tr>
<tr>
<td>Potassium sorbate</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
</tr>
</tbody>
</table>

Table 2. Results of Quality Control Testing of the Herbal Syrup

<table>
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<th>6</th>
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</thead>
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<td>Dark brown</td>
<td>Dark brown</td>
</tr>
<tr>
<td>Taste</td>
<td>Bitter-sweet</td>
<td>Bitter-sweet</td>
<td>Bitter-sweet</td>
</tr>
<tr>
<td>Appearance</td>
<td>Semi-clear liquid</td>
<td>Semi-clear liquid</td>
<td>Semi-clear liquid</td>
</tr>
<tr>
<td>Density (g/mL)</td>
<td>1.24 ± 0</td>
<td>1.24 ± 0</td>
<td>1.25 ± 0</td>
</tr>
<tr>
<td>Viscosity (cP)</td>
<td>144.4 ± 22.4</td>
<td>138.4 ± 15.4</td>
<td>286.4 ± 42.8</td>
</tr>
<tr>
<td>pH</td>
<td>5.39 ± 0.03</td>
<td>5.38 ± 0.03</td>
<td>5.34 ± 0.03</td>
</tr>
<tr>
<td>Dried residue (%)</td>
<td>47.82 ± 0.94</td>
<td>49.28 ± 0.87</td>
<td>49.58 ± 0.92</td>
</tr>
<tr>
<td>Total phenolics content (g/100 mL) (as pyrogallol)</td>
<td>0.434 ± 0.01</td>
<td>-</td>
<td>0.432 ± 0.02</td>
</tr>
</tbody>
</table>

* Data are reported as mean ± SD of the mean (n = 3).

herbal syrup and metformin compared to the PCOS group. The dose 2 group had the most reduction effect on ovaries weight.

4.3. Effects of the Herbal Syrup on Hormonal Levels

As shown in Figure 2, LH, FSH, LH/FSH ratio, estrogen, testosterone, and progesterone levels were measured in the serum of female rats. In the PCOS group, LH, LH/FSH ratio, and testosterone levels increased ($P < 0.001$), whereas estradiol and progesterone levels decreased ($P < 0.05$) compared to the control group. The reverse effect was observed in the metformin group as estradiol and progesterone levels increased. However, LH, LH/FSH ratio, and testosterone levels decreased ($P < 0.001$) compared to the PCOS group. In all doses of the herbal syrup groups, similar to the metformin groups, LH and testosterone levels decreased. LH levels significantly decreased in all herbal syrup groups, but testosterone reduction was prominent only in the dose 2 group ($P < 0.001$). In the herbal syrup groups, estradiol and progesterone increased to different levels in a dose-dependent manner. These increases in progesterone levels were not significant in the dose 1 group. There was no significant difference between all groups in FSH levels compared to the control group, and there was a significant reduction in the dose 2 group compared to the PCOS group ($P < 0.05$).

4.4. Effects of the Herbal Syrup on Histology of Ovaries

Many large ovarian cystic follicles and fewer corpora lutea were recognized in the PCOS group, whereas no histological abnormalities were observed in the control group. Histological studies of metformin and herbal syrup groups exhibited normal follicular development with a significant improvement in the number of primary, secondary, antral, and Graafian follicles, along with corpora lutea, compared to the PCOS group, which is associated with a significant decrease in the number of cystic follicles (Table 3 and Figure 3).

5. Discussion

PCOS seems to be a congenital disorder. It is also influenced by environmental factors that cause hormonal changes, including excess androgen production, with clinical features such as irregular menstrual periods, hirsutism, and acne (39). There is no definitive diagnostic test to recognize PCOS, and clinical diagnosis is based on the presence of 3 specific features, oligo-anovulation, androgen excess (either clinical or biochemical), and polycystic ovaries on ultrasound evaluation (40).

Since herbal medicine has long offered suitable therapies for women with irregular menstruation, hyperandrogenism, and PCOS (11), we decided to study the effect of
Figure 1. (A) Body weight difference before and after treatment. (B) Ovaries weight measurement. A 1-way analysis of variance followed by Tukey multiple comparisons. Data are expressed as mean ± SEM (n = 6 animals in each group). * P < 0.001 compared to the PCOS group, + P < 0.05 groups compared to the control group.

Figure 2. Mean ± SEM of LH, FSH, testosterone, progesterone, and estradiol concentrations in control, PCOS, metformin, and herbal syrup groups after 28 days of the experiment. (A) LH, (B) FSH, (C) LH/FSH ratio, (D) estradiol, (E) testosterone, and (F) progesterone. Values are expressed as mean ± SEM (n = 8). A 1-way analysis of variance followed by Tukey multiple comparisons. + P < 0.05; ++ P < 0.001 as compared to the control group.* P < 0.05; ** P < 0.001 as compared to the PCOS group.

herbal medicine to treat PCOS using ITM. A syrup formulation was prepared based on old prescriptions and current pharmaceutical standards. Fewer additives such as coloring, flavoring, and sweetening agents made the final syrup more similar to the traditional dosage form. In the original formulation of the syrup (28), fennel, anise, and celery seeds were mentioned as effective ingredients, and other excipients, such as sugar and CMC, were used to achieve the desired taste and viscosity.

As shown in Table 1, the F5 formulation with fewer extra ingredients, 10% plant materials, and the best taste and viscosity was chosen as the final preparation.

After 6 months of storage, the F5 samples were preserved their colors, tastes, and semi-clear appearance. Cap
Figure 3. Photomicrographs of the elected ovarian cross-section from control, PCOS, metformin, and all doses of herbal syrup treated rats after 28 days of the experiment. 
(A) The control group with different stages of ovarian follicles, including secondary, antral, Graafian follicles, and corpora lutea. 
(B) The PCOS group with various giant cystic follicles. 
(C) The metformin group with secondary, antral, Graafian follicles, and corpora lutea. 
(D) The dose 1 group with small cystic follicles and other follicles. 
(E) The dose 2 group with mature Graafian follicles, small cystic follicles, other developing follicles, and corpora lutea. 
(F) The dose 3 group with lots of secondary and antral follicles and small cystic follicles (Hematoxylin and eosin staining, x10).
Our findings demonstrated that the herbal syrup normal-
trogen levels induced by enzymatic inhibition of letrozole.

cumulation of androgens and a significant decrease in es-
crease in testosterone levels, presumably reflecting the ac-
perandrogenism. The PCOS group showed a significant in-
elss.
estrous cycles and an increase in testosterone and LH lev-
tration, PCOS induction was confirmed by irregularities in
be expected to result in increased ovarian androgen, de-
crease in the activity of this enzyme by letrozole could
androgens by aromatase produced by granulosa cells. A
(44). In ovaries, estradiol is generated by converting C19
sions, appropriate animal models that mimic many PCOS
model is similar to the human PCOS in many features such
the letrozole model has been increasing (42). The letrozole
and in recent years, the number of published studies on
PCOS was introduced in 2003, and in the next step, the therapeutic effect of the selected
formulation was evaluated on PCOS treatment through
the animal model. Due to the logistic and ethical limita-
tions, appropriate animal models that mimic many PCOS
characteristics have been developed since the 1960s (41).
The letrozole-induced-PCOS model was introduced in 2003, and in recent years, the number of published studies on
the letrozole model has been increasing (42). The letrozole
model is similar to the human PCOS in many features such
as hormonal imbalance, circulating hyperandrogenism,
and ovarian morphologic (43). Letrozole is an inhibitor
of the P450 aromatase enzyme that catalyzes the conver-
station of androgens (androstenedione and testosterone) to
estrogens (estrone and estradiol) during steroidogenesis
(44). In ovaries, estradiol is generated by converting C19
androgens by aromatase produced by granulosa cells. A
decrease in the activity of this enzyme by letrozole could
be expected to result in increased ovarian androgen, de-
creased estrogen, and development of PCOS in the animal
models (45). In our study, after 21 days of letrozole admin-
istration, PCOS induction was confirmed by irregularities in
estrous cycles and an increase in testosterone and LH lev-
els.

One of the main reproductive features of PCOS is hy-
perandrogenism. The PCOS group showed a significant in-
crease in testosterone levels, presumably reflecting the ac-
cumulation of androgens and a significant decrease in es-
trogen levels induced by enzymatic inhibition of letrozole.
Our findings demonstrated that the herbal syrup normal-
ized the hormonal level, and we found the best efficacy
at 2 and 4 mL/kg doses. These modifications are probably
due to phytoestrogen compounds (such as anethole in fenn-
el, safrole in anise, and apigenin in celery) and antian-
drogenic compounds (β-sitosterol and palmitic acid in 3
herbs). It has been revealed that β-sitosterol and palmitic
acid (found in fennel, anise, and celery) have antian-
drogenic properties and reduce testosterone by inhibiting the
dihydrotestosterone-receptor complex (46). Also, estro-
genic compounds reduce androgens by stimulating hep-
atic production of sex hormone-binding globulin (47).

In addition, one of the confirming indicators of PCOS is
elevated LH level (35). Excess androgen feedback to the
pituitary gland provokes excessive LH and decreases FSH
secretion (48). In this study, the herbal syrup was signifi-
cantly downregulated the LH levels compared to the PCOS
group. Besides the antiandrogenic compounds of 3 herbs,
chronic use of plant extracts containing phytoestrogens
lowers testosterone levels. In the herbal syrup group, prob-
ably because of the synergic effect of the 3 herbs, less LH
was produced due to the reduction of androgens. A simi-
lar study stated that fennel with this mechanism reduced
LH levels in PCOS rats (20). In PCOS, the pulse frequency of
gonadotropin-releasing hormone (GnRH) increases; there-
fore, the FSH release decreases in a more significant pro-
portion to LH, and the LH/FSH ratio increases (49). The
LH/FSH ratio is typically 1: 1; however, this ratio is 2 or
3 times higher (2: 1 or 3: 1) in PCOS cases (18). A re-
cent study shows that normalizing the LH/FSH ratio is es-
sential in PCOS treatment (50). Furthermore, our results showed
that the LH/FSH ratio in PCOS rats was 2.3 times higher
than in the control group. This ratio became normal af-
fter the herbal syrup administration. Our study does not
change FSH levels significantly, similar to previous studies
(51, 52). Thus apparently, a decrease in LH levels reduces
the LH/FSH ratio. In PCOS women, progesterone blood levels
decrease due to anovulation and reduction in the number

| Table 3. Mean Follicle Numbers in Different Groups (Mean ± SEM) |
|-------------------|-------------------|-------------------|-------------------|-------------------|-------------------|
|                   | Primary Follicle  | Secondary Follicle | Antral Follicle   | Graafian Follicle | Cystic Follicle   |
| Control           | 6.24 ± 1.17       | 6.3 ± 0.59        | 3.18 ± 0.89      | 1.60 ± 0.80       | 0 ± 0              |
| PCOS              | 0.50 ± 0.24       | 3.29 ± 0.33       | 1.47 ± 0.53      | 2.24 ± 0.95       | 13.68 ± 5.84      |
| Metformin         | 3.25 ± 0.57       | 5.47 ± 0.88       | 3.69 ± 0.74      | 0.44 ± 0.72       | 2.53 ± 0.25       |
| Dose 1            | 4.04 ± 0.53       | 4.44 ± 0.64       | 3.03 ± 0.65      | 0.91 ± 2.57       | 2.39 ± 2.57       |
| Dose 2            | 4.19 ± 0.46       | 6.44 ± 0.91       | 4.19 ± 0.69      | 1.11 ± 2.57       | 3.69 ± 2.57       |
| Dose 3            | 3.16 ± 0.75       | 8.25 ± 1.62       | 6.08 ± 1.77      | 0.66 ± 2.57       | 3.36 ± 2.57       |

a P < 0.001 as compared to the control group.
b P < 0.05
c P < 0.001 as compared to the PCOS group
d P < 0.01
of corpora lutea (53), and our results agree with these findings. The herbal syrup increased the number of corpora lutea due to lowering LH levels, improving the process of folliculogenesis, and increasing ovulation rates. The corpora lutea secrete the most progesterone and ultimately increase progesterone blood levels. Previous studies have investigated the effects of fennel on improving the hormonal profiles in PCOS and other female disorders, which are consistent with our results (20, 54). Furthermore, the combination of anise and celery regulated menstrual cycles and improved oligomenorrhea in PCOS patients (15). As a standard treatment, metformin improved blood levels of sex hormones significantly, according to the results of the previous study (35).

Obesity is one of the complications that affect PCOS women, with a worldwide prevalence of approximately 35% to 80% (55). The letrozole model was exhibited weight gain in PCOS rats, aligning with our results (43). As an ingredient of this syrup, celery contains a flavonoid named luteolin. Evidence suggests that luteolin suppresses mast cell secretion, especially interleukin (IL)-6, regulates connected adipose tissue angiogenesis, reduces body weight, reduces inflammation in adipose tissue, and lowers insulin resistance (56). The significant weight loss seen in the herbal syrup groups may be due to this combination of celery.

In line with previous studies, our findings indicate that ovarian weight was increased in the PCOS group (48, 57). The herbal syrup and metformin could prevent ovarian weight gain induced by PCOS. This weight loss can be due to a reduction in the number and size of cystic follicles and an improvement in the folliculogenesis process. In PCOS women, hyperandrogenism and elevated LH levels cause multiple cysts to form in the ovaries (58). While LH levels increase, androgens production in the theca cells is accelerated. Subsequently, the relative FSH deficiency reduces the conversion of androgen into estrogen by granulosa cells and disrupts follicle maturation and ovulation (59). In our study, after induction of PCOS, excess androgen production resulted in growth disturbance of antral and Graafian follicles (60). Therefore, the number of different follicles decreased alongside sizeable cystic follicles with a thin granulosa layer. Baravalle et al. showed that letrozole generated follicular dysfunction, such as atretic and large cysts with a thin granulosa layer (61). In our study, the herbal syrup reduced the number of cysts and increased the thickness of the granulosa layer, demonstrating the improvement of ovarian tissue in these groups. The herbal syrup improved folliculogenesis and increased the number of primary, secondary, antral, and Graafian follicles. These increases have also been reported after fennel administration due to estrogenic properties (62). Besides, the antioxidant features of fennel, anise, and celery could effectively prevent ovarian follicle atresia by diminishing oxidative stress (61, 63).

5.1. Conclusions

Our study exhibited that the herbal syrup (containing fennel, anise, and celery) could recover ovarian morphology and upregulate serum levels of progesterone and testosterone. In contrast, it could decrease LH, estradiol, and FSH serum levels in the letrozole-induced PCOS model. The studied herbal syrup might be a potent therapeutic herbal drug for PCOS treatment. Further clinical investigations are required to confirm the potential therapeutic effects of the recently studied herbal syrup.

Footnotes

Authors’ Contribution: Sh. F., Sh. S., and Z. Kh. contributed to the development of the protocol, abstracted data, performed the experiments, and prepared the manuscript. M. T. participated in designing the evaluation. Z. Sh. contributed to the development of the protocol. M. F. re-analyzed the clinical and statistical data and revised the manuscript. All authors read and approved the final manuscript.

Conflict of Interests: The authors declare no conflicts of interest and are responsible for the content presented in the paper.

Data Reproducibility: The data presented in this study are uploaded during submission as a supplementary file and are openly available for readers upon request.

Ethical Approval: The Ethics Committee of Shahid Beheshti University of Medical Sciences approved this study (code: IR.SBMU.RETECH.REC.1399.899, 2021). The in vivo step was done according to the NIH Animal Care and Use Committee Guide for the Care and Use of Laboratory Animals.

Funding/Support: This project was granted by Traditional Medicine and Materia Medica Research Center, Shahid Beheshti University of Medical Sciences, Tehran, Iran (grant number: 188/96). The institute had no role in the design of the study, collection, analysis, interpretation of data, or writing the manuscript.

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