Tanshinone IIA attenuates estradiol-induced polycystic ovarian syndrome in mice by ameliorating FSHR expression in the ovary

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Abstract. Tanshinone IIA (TSIIA) is a major component of *Salvia miltiorrhiza*, a Chinese herb that exhibits a therapeutic effect on polycystic ovary syndrome (PCOS). The present study replicated PCOS via the neonatal treatment of estradiol in mice. Estrous cycles, body and ovarian weight, serum levels of testosterone and estradiol were determined. Histological examination of ovaries was performed. The mRNA and protein levels of aromatase luteinizing hormone receptor and follicle-stimulating hormone (FSHR) in ovaries and granule cells were assayed by reverse transcription-quantitative polymerase chain reaction and western blotting, respectively. TSIIA was revealed to reverse all disorders induced by estradiol treatment, including prolonged estrous cycles, increased body and ovarian weight, increased atretic cyst-like follicles and decreased corpus luteum, large antral follicles and preovulatory follicles. These improvements in PCOS as a result of TSIIA treatment are likely due to the revised testosterone/estradiol balance, as TSIIA reversed the decrease in aromatase mRNA, the enzyme that converts androgen to estrogen. As the expression of aromatase is regulated by the FSH pathway, TSIIA-mediated elevation in FSHR expression may lead to the upregulation of aromatase. Therefore, TSIIA revises the balance of androgen and estrogen by rescuing the reduced expression of FSHR and aromatase, thus attenuating murine PCOS. The current study aimed to further the application of natural drugs in the treatment of PCOS to confront the side effects of hormone drugs and expand the use of TSIIA.

Introduction

Polycystic ovarian syndrome (PCOS) is the most common endocrine and metabolic disease in women, occurring in ~15% at procreation age (1). Its characteristics include hyperandrogenism, ovulation failure and polycystic ovarian morphology (2), combined with multiple clinical symptoms including menoxenia, acne, hirsutism, alopecia, central obesity and infertility (3). PCOS, is a life-long disease, which not only causes short-term problems, including infertility due to anovulation, but also exhibits long-term metabolic issues including obesity, insulin resistance, non-alcoholic fatty liver disease, type 2 diabetes mellitus and cardiac vascular disease (4), as well as hyperinsulinemia and estrogen dependent cancer (5).

Hyperandrogenism, occasionally combined with hypoestrogenism, is a classic diagnostic criterion for PCOS, as excess androgen induces the dysfunction of the hypothalamic-pituitary-ovarian axis, which is the neurohormonal system that regulates gonadal hormone balance, follicle maturation and ovulation (6). Mainstream therapy including hormonal therapy, antiandrogens and insulin-sensitizing drugs used to reduce androgen levels is usually combined with oral contraceptives (7). Among them, diane-35 (a combination of ethinyl estradiol and cyproterone acetate) is commonly used to treat PCOS (8).

However, diane-35 produces significant hormonal side effects and negatively impacts triglyceride metabolism and pancreatic function (9).

*Salvia miltiorrhiza* (*S. miltiorrhiza*), a Chinese herbal medicine, has been used to treat PCOS. Components isolated from *S. miltiorrhiza*, including cryptotanshinone, attenuate PCOS by reversing dexamethasone-induced ovarian insulin resistance in mice (10). However, whether other components of *S. miltiorrhiza* have the capability to treat PCOS remains unclear. Tanshinone IIA (TSIIA) is another primary component of *S. miltiorrhiza* (11). Serving as an antioxidant, anti-inflammatory and immunomodulatory element, it is primarily used to treat cardiovascular diseases (12). TSIIA also inhibits oxidative stress injury in neuronal cells to treat Alzheimer's disease (13). The results of the present study confirmed the therapeutic effect of TSIIA on PCOS induced by neonatal...
exposure to estradiol in mice by regulating follicle-stimulating hormone (FSH) receptor (FSHR) expression in the ovary.

**Materials and methods**

**Animals.** Animal care and experiment procedures were performed in accordance with the guidelines of and approved by the Animal Care and Ethical Committee of Affiliated Hospital of Nanjing University of Chinese Medicine (Nanjing, China). A total of 57 female (C57BI/6J x A/J) F1 (B6A) 5-day-old mice (weight, 4-5 g) purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd. (Beijing, China) were used in the current study. All mice were maintained in standard cages under a 12 h light/dark cycle at a temperature of 22-24°C and a relative humidity of 55-65%, with ad libitum access to food and water.

**Experimental design and drug administration.** Estradiol (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) dissolved in dimethyl sulfoxide and diluted with sesame oil (BSZH Co. Ltd., Beijing, China), was subcutaneously injected into 5-day-old female mice (PCOS mice, n=56) at a dose of 20 µg/day for 3 days (14). Vehicle of the same volume was administrated to mice (control group, n=28). On the 56th postnatal day, half PCOS mice (estradiol+TSIIA group, n=28) received TSIIA (Nanjing Spring and Autumn Biological Engineering Co. Ltd, Nanjing, China) for 4 weeks at a dose of 100 mg/kg/day by gavage. The PCOS mice left (estradiol group, n=28) and control mice received the same weight-based vehicle. Estrous cyclicity was determined during the last 18 days prior to sacrifice. Thereafter, mice were weighed (22-25 g) and sacrificed on the 96th postnatal day by decapitation.

**Assessment of estrous cycles.** Estrous cycles were examined daily at 0800-0900 h. The fluid obtained by vaginal lavage with 0.9% saline was spotted thinly on a microscope slide. Following air-drying, slides were stained with toluidine blue (0.1%) at room temperature for 2 sec. According to the types of vaginal epithelial cells present (leukocyte, nucleated and cornified cells), diestrus, estrus and proestrus were identified using light microscopy at magnification, x100 as previously described (15).

**Ovarian histology.** Ovaries were excised from mice, fixed in 4% paraformaldehyde at room temperature for 48 h, dehydrated in ascending grades of ethanol and embedded in paraffin in 5-µm sections. Samples were then deparaffinized, rehydrated and stained with hematoxylin and eosin (H&E) for 3 and 0.5 min at room temperature, respectively. Follicles were counted using a conventional light microscope (Olympus DP70; Olympus Corporation, Tokyo, Japan) with x40 objective. The number of follicles (atretic cyst-like, large antral and preovulatory follicles) were counted in every sixth section (30 µm apart) and multiplied by 6 to provide the total number of follicles in each ovary. Only follicles containing an oocyte with a visible nucleus were counted to avoid double counting. The classification of follicular stages was made following the morphological criteria as described previously (16,17). The number of corpora lutea was scored in a blinded fashion using one section per ovary and one ovary per mouse (18).

**Hormonal measurements.** Serum was obtained when mice were sacrificed following the determination of estrous cyclicity. Levels of FSH, luteinizing hormone (LH), progesterone (P), estradiol and testosterone (T) were quantified using ELISA kits (cat. nos. E0830Mu, E0441Mu, E0459Mu, E0461Mu and E0458Mu, respectively; UsCN Life Sciences, Inc., Wuhan, China) according to the manufacturer’s protocol.

**Reverse transcription quantitative polymerase chain reaction (RT-qPCR).** Total RNA was extracted from murine ovary tissue using a TRIzol reagent kit (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA) according to the manufacturer’s protocol. RNA (1 µg) was transcribed into cDNA using a PrimeScript™ RT reagent kit (TaKaRa Biotechnology Co., Ltd., Dalian, China). qPCR reactions were performed using a Light Cycler Fast Start DNA Master SYBR Green I kit and an ABI Prism 7300 Sequence Detection System (Applied Biosystems; Thermo Fisher Scientific, Inc.) The thermocycling conditions were as follows: 95°C for 3 min; 95°C for 15 sec; 60°C for 15 sec; 72°C for 1 min (35 cycles); and 72°C for 10 min. The relative expression of genes was determined using the 2−ΔΔcq method (19) with normalization to 36B4 expression. The primer sequences were as follows: LH receptor (LHR) forward, 5'-ATGAGTCCATCAAGTCTGAAAC-3' and reverse, 5'-CCTGCAATTTGGTGGAAAGAGA-3'; FSHR forward, 5'-CTTGCTCTTGTGTCCTCCTTG-3' and reverse, 5'-CTCGGTACCTGTGTTACATCTTTG-3'; aromatase forward, 5'-ATGGTTGGAATGCTGAACCCC-3' and reverse, 5'-AGGACCTGTATTTGAGACCGAG-3'; peroxisome proliferator-activated receptor γ (PPARγ) forward, 5'-TTTCTCCAGAAGAACCATCGATT3'-3' and reverse, 5'-ATGGCAATTCTGGAGACATCCTCC-3'; and 34B4 forward, 5'-AGGGCGCTCTCTGGCATCT-3' and reverse, 5'-CCCGAGGGGCAGATCTTGG-3'.

**Isolation of granule cells.** As described previously (20), ovaries were isolated from mice and the large and transparent follicles residing within were punctured to liberate granule cells into the medium, Dulbecco's modified Eagle's medium (DMEM)/nutrient mixture F-12 (F12; Gibco; Thermo Fisher Scientific, Inc.). Granule cells were mixed and collected via centrifugation at 100 x g at room temperature for 2 min. Cells were then suspended in DMEM/F12 complete medium containing 10% fetal bovine serum (Gibco; Thermo Fisher Scientific, Inc.) and 1% nonidet P-40, 10 mM NaF, 1 mM NaVO4, 1 mM phenylmethylsulfonyl fluoride, 10 mg/ml aprotinin, 10 mg/ml leupeptin; all components purchased from Aladdin Shanghai Biochemical Technology Co., Ltd., Shanghai, China) and boiled at 95°C for 5 min in 5X loading buffer.
[250 mM Tris-HCl (pH 6.8), 10% SDS, 0.5% bromophenol blue, 50% glycerol, 5% mercaptoethanol; all components purchased from Aladdin Shanghai Biochemical Technology Co., Ltd.]. Protein concentration was assessed using a bicinchoninic acid protein assay kit (Pierce; Thermo Fisher Scientific, Inc.). Protein (25 µg) was separated by 10% SDS-PAGE and transferred to polyvinylidene difluoride membranes. The membranes were incubated with rabbit anti-FSHR antibodies (1:1,000; cat. no. K002799P; Beijing Solarbio Science and Technology Co., Ltd., Beijing, China), anti-LHR antibodies (1:1,000; cat. no. sc-25828; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), anti-aromatase antibodies (1:1,000; cat. no. PA1-21398; Invitrogen; Thermo Fisher Scientific, Inc.) and mouse anti-β-actin (1:1,000; cat. no. CBL171-I; EMD Millipore, Billerica, MA, USA) at 4˚C overnight. Membranes were then further incubated with horseradish peroxidase-conjugated goat anti-rabbit Immunoglobulin G (IgG; 1:5,000; cat. no. AP183P) or goat anti-mouse IgG (1:5,000; cat. no. AP192P; both EMD Millipore) for 2 h at room temperature. Membranes were visualized using the Tanon-5200 Chemiluminescence Imager (Tanon Science and Technology, Co., Ltd., Shanghai, China) with an ECL Western blotting substrate (EMD Millipore). ImageJ version 1.48 (National Institutes of Health, Bethesda, MA, USA) was used for the densitometry analysis of bands.

Cyclic adenosine monophosphate (cAMP) levels in granule cells. Following culture for 24 h, non-adherent granule cells were removed and adherent cells were treated with FSH (100 IU/l, Ningbo Second Hormone Factory, Ningbo, China) for 10 min at 37°C. Cells were then rinsed with cold PBS once and lysed using 0.1 M hydrochloric acid for the protein concentration assay with total protein assay kit (EMD Millipore). ImageJ version 1.48 (National Institutes of Health, Bethesda, MA, USA) was used for the densitometry analysis of bands.

Statistical analysis. Data were expressed as the mean ± standard error of the mean. Data between groups were analyzed using one-way analysis of variance followed by Dunn’s multiple comparisons test. P<0.05 was considered to indicate a statistically significant result.

Results

TSIIA attenuates estrous cycle disorder in mice with PCOS. Normal control mice exhibited a regular estrous cycle of 5 days on average (Fig. 1A). The entire estrous cycle comprises four phases, which includes proestrus, estrus, metestrus and diestrus. Neonatal estradiol-treatment increased metestrus and diestrus compared with control mice, which contributed to the prolongation of the estrous cycle to ~9 days (Fig. 1A and B). TSIIA treatment significantly reduced entire estrous cycle, metestrus and diestrus and led to a slight increase in the estrous period (Fig. 1A and B).

Effects of TSIIA on ovarian morphology in mice with PCOS. Body weight and ovary weight in each group were assayed following sacrifice. The results revealed that neonatal estradiol treatment increased body and ovarian weight in mice (Fig. 2A and B). The increase in ovarian weight was more pronounced than that of body weight, and the ratio of ovary to body weight also significantly increased following estradiol treatment (Fig. 2C). TSIIA treatment reversed the estradiol-induced increase in body weight, ovary weight and the ratio of ovary to body weight (Fig. 2A-C).

Pathological changes in murine ovaries were assessed via H&E staining. In congruence with a previous study (14), multiple enlarged follicles were observed in the ovaries of estradiol-treated mice in the present study (Fig. 2D), which mimicked PCOS. Neonatal estradiol-treatment reduced the number of large antral follicles, preovulatory follicles and corpus luteum, but increased the number of atretic cyst-like follicles (Fig. 2D), which were also similar to changes typically observed in PCOS (6). TSIIA treatment reversed the observed pathological change in the number of follicles (Fig. 2D).
TSIIA regulates serum hormone levels in mice with PCOS. Ovarian structure and function is primarily regulated by serum hormones (6). Thus, levels of FSH, LH, estradiol, T and P in murine serum were assayed. As presented in Fig. 3A-C, serum levels of LH, FSH and the ratio of LH to FSH were not significantly affected by neonatal estradiol-treatment nor by TSIIA. However, serum estradiol was reduced and T levels were elevated following neonatal estradiol treatment, which were each reversed when further treated with TSIIA (Fig. 3D and E). Serum P levels were also reduced following neonatal estradiol-treatment, but TSIIA demonstrated no significant effect.

Effect of TSIIA on LHR, FSHR, aromatase and PPARγ expression in the ovaries of mice with PCOS. Neonatal mice treated with estradiol exhibited significantly lower mRNA levels of aromatase, the key enzyme for the transferal of T to estradiol in ovaries (21). However, mRNA levels of ovarian aromatase recovered following TSIIA treatment. FSHR, LHR and PPARγ function to regulate the expression of aromatase in the ovary (22-24); however, only FSHR exhibited significantly lower mRNA levels following neonatal estradiol treatment, which was recovered by TSIIA (Fig. 4A).

In ovarian granule cells (Fig. 4B-D) of neonatal estradiol-treatment mice, western blot analysis revealed lower FSHR and aromatase protein levels, which were improved following TSIIA treatment (Fig. 4B).

The FSH-induced cAMP levels in granule cells isolated from estradiol-treated mice were significantly lower than that from control mice, whereas TSIIA treatment elevated the granular cAMP level in estradiol-treated group similar to control group (Fig. 4C).

The results also demonstrated that granular FSHR and aromatase mRNA levels in estradiol treated group were significantly lower than in the control group, but were reversed following TSIIA treatment (Fig. 4D).

Discussion

The present study revealed the therapeutic effect of TSIIA, a component isolated from Salvia miltiorrhiza, on murine PCOS induced by the neonatal treatment of estradiol.

In a previous study (25), PCOS was successfully replicated in mice via neonatal treatment with estradiol. This was determined as mice exhibited a prolonged diestrus phase,
enlarged ovaries, a thickened theca layer, a decreased number of corpus luteums and preovulatory follicles, an increased number of atretic follicles and multiple cysts in ovaries, which are characteristics similar to those in women with PCOS (19). In the current study, estrus cycles were returned to normal status following TSIIA treatment and this also reversed the pathological change of murine ovaries with estradiol-induced PCOS.

The alteration of granulosa cell hormonal equilibrium is a cause of disturbed follicular development and maturation that results in the premature arrest of follicular growth in PCOS (26,27). Estrogen suppresses granulosa and luteal cell apoptosis (28,29) and triggers the release of FSH and LH, promoting follicle maturation and ovulation (30). An excess of intraovarian androgen leads to follicular atresia (28), as it activates primordial follicles via the phosphatidylinositol-3-kinase/protein kinase B/Forkhead box O3a pathway and inhibits growth differentiation factor 9 expression in oocytes, resulting in the arrest of preantral follicle development (31).

In the present study, neonatal treatment of estradiol decreased serum estradiol but increased serum T levels, which may lead to pathological and functional changes in the ovary. Thus, the therapeutic effect of TSIIA observed in the current study may be the result of disordered estradiol and T, as other hormones affecting follicle maturation, including LH, FSH and P, were either not changed or not reversed following TSIIA therapy.

The hormonal balance between estrogen and androgen is primarily sustained by the normal function of granulosa cells, which contain a series of enzymes converting T to estradiol. Disorder in the activity or expression of aromatase, the key enzyme catalyzing the conversion of T to estradiol, induces excessive androgen accumulation in serum (32). Thus, aromatase mRNA levels of the ovaries and granule cells were determined in the current study. It was demonstrated that they were reduced by neonatal estradiol-treatment, but ameliorated following TSIIA therapy. This implies that TSIIA targets the expression of aromatase to revise the T/estradiol balance in mice.

The transcription of aromatase is regulated by the LH, FSH and PPARγ pathways (22-24). LH and FSH mediate the down- or upregulation of aromatase via LHR and FSHR, respectively. Since the serum levels of LH and FSH were not affected by neonatal estradiol treatment or TSIIA in the current study, the expression of LHR and FSHR should predict the transcription of aromatase. FSHR mRNA levels were reduced by neonatal estradiol treatment, which was then ameliorated by TSIIA. However, neither LHR nor PPARγ expression was affected. The results indicate that the downregulation of FSHR induced PCOS in the present murine model and that TSIIA exerted its therapeutic effect by targeting FSHR expression. Furthermore, the present study revealed that estradiol treatment reduced FSHR and aromatase protein levels, thus inhibiting FSH-induced cAMP elevation in granule cells isolated from mice. TSIIA treatment also ameliorated FSHR expression and
an FSH induced cAMP increase. As a result, TSIIA primarily affected the ovarian expression of FSHR, which improved ovarian function.

In summary, the present study revealed that PCOS induced by neonatal exposure to estradiol in mice is attenuated by TSIIA treatment. The therapeutic effect of TSIIA is likely due to the recovery of FSHR expression in the ovary, thus leading to elevated serum levels of estradiol and T, an improvement in the structure of ovaries and the attenuation of the disordered estrous cycle. The current study revealed a novel monomer (TSIIA), which may be utilized to further the application of natural drugs in the treatment of PCOS to reduce the side effects of hormone drugs. However, the present study was limited by the lack of a positive control group and intensive studies in granule cells.

Figure 4. Effect of TSIIA on the mRNA expression of genes associated with ovarian hormone levels. (A) mRNA levels of LHR, FSHR, aromatase and PPARγ in ovaries (n=8 for each group). Granule cells were isolated from Estradiol, Estradiol+TSIIA and control mice. (B) FSHR, LHR and aromatase protein levels in granule cells. (C) cAMP levels in granule cells treated with/without FSH. (D) mRNA levels of FSHR and aromatase in granule cells. Data are expressed as the mean ± standard error of the mean. *P<0.05. TSIIA, Tanshinone IIA; LHR, luteinizing hormone receptor; FSHR, follicle stimulating hormone receptor; PPARγ, peroxisome proliferator-activated receptor γ; cAMP, cyclic adenosine monophosphate.
Further study is therefore required to assess the mechanism of how TSIIA upregulates FSHR in granule cells.

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Availability of data and materials

All data and materials supported the results of the present study are available in the published article.

Authors' contributions

HFZ was responsible for the experimental design and drafting the manuscript. JJ, QYH, WWX, WJZ, BL, JL and WW performed the experiments and data analysis. HFZ reviewed the manuscript. All authors gave final approval of the version to be published.

Ethics approval and consent to participate

Animal care and experimental procedures were approved by and performed in accordance with the guidelines of the Animal Care and Ethics Committee of Affiliated Hospital of Nanjing University of Chinese Medicine.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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