Troxerutin protects against DHT-induced polycystic ovary syndrome in rats

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**Abstract**

The exact pathogenesis of polycystic ovary syndrome (PCOS), the most common neuroendocrine disorder in women of reproductive age, has not been fully elucidated. Recent studies suggested that chronic inflammation and neurotransmitter disorder involved in the progress of PCOS. Troxerutin, a natural flavonoid, was reported to possess neuroprotective effect in several disease models by inhibiting inflammation or enhancing neurotrophic factor. In this study, we investigated the possible protective effect and mechanism of troxerutin in a dihydrotestosterone (DHT)-induced rat model of PCOS. The PCOS rat models were treated with troxerutin at a dose of 150mg/kg or 300mg/kg for up to 4 weeks. Results showed that 300mg/kg troxerutin significantly decreased the body weight gain and improved the pathological changes of ovary induced by DHT. Meanwhile, the elevated gonadotrophin-releasing hormone (GnRH), gonadotrophin and testosterone in the serum of PCOS rats were reduced with the treatment of troxerutin. The expression of kisspeptin and NKB in arcuate nucleus and their receptors kiss1r and NK3r in GnRH positive neurons of median eminence were markedly decreased in troxerutin-treated rats. Of note, the GnRH inhibitory regulator GABA and stimulatory regulator glutamate were also restored to the normal level by troxerutin. The present study indicated that troxerutin may exhibit a protective effect in PCOS rat model via regulating neurotransmitter release.

**Introduction**

Polycystic ovary syndrome (PCOS) is a reproductive endocrinopathy with the prevalence estimated to be 6% ~ 20% (depending on the different diagnostic criteria used), making it the most common endocrine condition in women of reproductive age [1]. The diagnostic features of PCOS include androgen excess, ovulatory dysfunction and polycystic ovaries[1]. Obesity is present in 30-60% of PCOS patients, depending on the country of origin[2, 3]. The most consistent biochemical abnormality in women with PCOS is hyper-secretion of androgen, elevated serum luteinizing hormone (LH) levels and low to normal serum follicle stimulating hormone (FSH) levels[4, 5]. Animal models that reflect PCOS features are crucial resources to investigate this syndrome. A chronically 5α-dihydrotestosterone (DHT)-treated rat model closely mimics the human PCOS phenotype and is a suitable model for investigations about PCOS.

The hypothalamus-pituitary-gonad (HPG) axis plays a central role in the exquisite neuroendocrine regulation of reproduction. Hypothalamic secretion of gonadotrophin-releasing hormone (GnRH), has been robustly established as the key pathway that controls reproductive function[6]. Axons of GnRH neurons protrude into the median eminence and release GnRH, which via hypophyseal portal system, enters anterior pituitary. Anterior pituitary, the primary target of GnRH, responds to stimulation by increasing the secretion of LH and FSH, which in turn lead to steroid production from the ovaries and stimulate follicle genesis and ovulation[7]. Thus, GnRH is known as the master hormone and provides the final common output of the network that regulates reproductive function.
GnRH is influenced by extrinsic factors such as neurotransmitters and neuropeptides, any alteration of the GnRH regulatory neurotransmitters, such as inhibitory gamma-aminobutyric acid (GABA) and stimulatory glutamate (Glu), may result in reproductive endocrine dysfunction[8]. Actually, the chronic anovulation characteristic of PCOS is attributed to increased central GnRH drive and resulting gonadotrophin aberrations, which likely result from the cumulative effect of altered GnRH stimulatory and inhibitory neurotransmitter in hypothalamus and pituitary gland[9]. Kisspeptin and subsequently discovered neurkinin B (NKB), two novel GnRH regulatory neuropeptides proved to be essential for normal GnRH secretion in humans, have come under intense spotlight in the last decade[10, 11]. The related discovery of kisspeptin-/-neurkinin B-/-dynorphin-(KNDy) pathway has further strengthened the understanding of GnRH secretion modulation[12]. KNDy neurons residing in the arcuate nucleus region of rodents co-express kisspeptin, NKB and DYN[12, 13]. NKB and kisspeptin derived from KNDy stimulate the release of GnRH by binding Kisspeptin 1 receptor (Kiss1r) and neurokinin 3 receptor (NK3r) expressed on GnRH neurons[14, 15].

Troxerutin, a trihydroxyethylated derivative of the naturally occurring plant bioflavonoid rutin, also called vitamin P4. It has been widely identified in plants consumed as part of our daily diet, such as tea, coffee, cereals, and a variety of fruits and vegetables, and exerts multifarious biological activities, such as antioxidant and anti-inflammatory effects and exerts beneficial effects on animal models of different central nervous system diseases [16, 17]. Troxertuin attenuated cognitive deficit and oxidative stress in mouse brain through decreasing reactive oxygen species[18], inhibited hippocampal neuron apoptosis in a rat model of Alzheimer's disease[19]. Troxerutin inhibited cyclin-dependent kinase 1 expression, enhanced type 1 protein phosphatase α dephosphorylation and abolished MEK/ERK1/2/C/EBP β activation, which subsequently reversed the memory impairment in the domoic acid-treated mice[20], and exhibited neuroprotective potential in 6-OHDA rat model of Parkinson's disease through mitigation of apoptosis, astrogliosis and oxidative stress [21], prevented the harmful effects of maternal high-fat diet on their offspring through inhibition of pro-inflammatory cytokines and elevation of brain-derived neurotrophic factor levels[22], significantly promoted nerve growth factor (NGF) mRNA expression and attenuated cognitive impairment and brain oxidative stress through the activation of NGF/TrkA signaling pathway in D-gal-injected mice[23]. However, there was no evidence that troxerutin had any effect on IGF-1 so far. Since the hypothalamus plays an important role in reproduction, we supposed that troxerutin may have a protective effect against reproductive neuroendocrine dysfunction in PCOS. This study aimed to address this issue and preliminarily investigate the potential mechanism underlying its action.

**Materials And Methods**

**Animals and troxerutin administration**

Postnatal 14 days female SD rats were purchased from the Breeder center of Rodents (Jinan, China). The rats were housed under controlled lighting (12h light, 12h darkness, lights were turned on at 08:00 and turned off at 20:00) and temperature (24°C) with free access to food and water for a week of acclimatization. All experimental protocols used in this study were approved by the Animal Care and
Ethical Committee of Xuzhou medical University. Postnatal 21 days rats weighing approximately 50-55g, were randomly divided into four groups: sham group (each rat was implanted with a tube without DHT), PCOS group (each rat was implanted with a silastic tube filled with 5α-DHT at dose of 7.5mg under anesthesia), PCOS+150mg/kg troxerutin group and PCOS+300mg/kg troxerutin group (PCOS rat with intraperitoneal injection of 150mg/kg or 300mg/kg troxerutin daily during the 4 weeks after implantation). Due to the high water-solubility of troxerutin, the gastrointestinal system can absorb it well without toxicity. Troxerutin interacts with tRNA by external binding manner with low binding affinity, and also binds with DNA through groove binding manner. Sham and PCOS groups were injected daily with equivalent 0.9% saline which was used to dissolve troxerutin. DHT was procured from sigma, St. Louis, Mo and troxerutin (purity >95%) was from Baoji Fangsheng Biotechnology Co., Ltd, Baoji, China. The rats were housed individually after surgery.

**Blood and tissue sampling**

At the terminal of the study, rats were decapitated and trunk blood was collected and centrifuged at 3000 rpm for 15 minutes. Plasma was separated and stored at -80°C until analyzed for biochemical and hormonal analysis. Ovaries were cleaned in saline and managed fat free. Ovaries were fixed in 10% buffered formalin for 48 hours and paraffin-embedded. Paraffin-embedded tissue sections were de-waxed, sectioned (6 μm) and stained with hematoxylin and eosin (H&E). Numbers of follicles were classified and counted in every 40th ovarian section for 10 ovaries per group. Follicles were counted according to the following definition. Primordial follicle was surrounded by a single layer of squamous granulosa cells. Primary follicle contained a single layer of cuboidal granulosa cells. Secondary follicle contained two or more layers of granulosal cells, but no antral space in the granulosa layer. Antral follicle displayed multi-layers of granulosal cells with the presence of follicular antrum.

**Vagina smears and estrous cycles**

Vaginal smears were taken daily at 09:00am from the 18th to the 28th day after the first day of DHT treatment, which were the terminal ten days of the study. Microscopic analysis (BX41, Olympus) was carried out to determine the stage of the estrous cycle by the predominant cell types in vaginal smears following methylene blue staining (Solarbio, Beijing). Proestrus consists of enlarged, round, nucleated epithelial cells; estrus consists of a large number of cornified squamous epithelial cells; metestrus consists of two kinds of cells: epithelial cells and leukocytes with approximately same amount; and diestrus is characterized by a small amount of cells with the predominance of leukocytes.

**Body weight, biochemical and western blot analysis**

Body weight was measured every week after implantation for a total of 4 weeks.

The serum concentration of LH, FSH and testosterone were measured via Enzyme Linked Immuno Sorbent Assay (ELISA) with the help of commercial kits (ELISA kit, CUSABIO, Inc, Wuhan, China) and the
procedure was followed as given in the kit catalog. The concentration of LH, FSH and testosterone was estimated by the standard curve.

To identify the hypothalamic GnRH status, hypothalamus was dissected out. The rat brain was carefully taken out and temporarily put on ice with the ventral surface upward. Thalamus is the ellipsoidal part at the center of the ventral brain. With the middle of tuber cinereum and optic chiasma as the center, prechiasmal border as the anterior edge, postmammillary border as the posterior edge, hypothalamus (4x4x2mm) was carefully dissected out. Hypothalamus tissue were homogenized to obtain protein samples. Then the protein (25μg) was separated by SDS-PAGE and transferred to membranes. The membranes were incubated with mouse monoclonal anti-GnRH antibody (1:1000, MAB5456-C, Millipore, Billerica, MA) and mouse anti-GAPDH antibody (1:20000, Proteintech, Chicago, USA). After being washed with TBST three times, the membranes were incubated with IRDye-labeled secondary antibodies in TBST for 2h. The bands on the membrane were scanned with an Odyssey infrared scanner (LI-COR Biosciences, Lincoln, NE, USA) and the density of the bands was analyzed with ImageJ software.

Chromatographic analysis

Chromatographic analysis was carried out to evaluate the levels of two kinds of GnRH regulatory neurotransmitters GABA and glutamate in the brain/pituitary. Weigh the brain and add in the mix of methanol and water (1:1) at the ratio of 100mg/ml to prepare homogenate. The brain homogenate was centrifuged at 12000 rpm for 30 min at 4°C. The mix of supernate and 80μL acetonitrile were centrifuged at 12000 rpm for 30 min at 4°C. The new supernate was derivatized, mixed with 80μL borate saline buffer, 160μL water and 80μL F-moc, filtered by 0.45μm filtration membrane and immersed in 40°C bath for 5 minutes. Then, the samples were used for the measurement of GABA and glutamate using chromatographic analysis. All standard biomarkers used for identification purpose in chromatographic studies were procured from Zhongke Co, Ltd, Beijing, China. High-performance liquid chromatography (HPLC) was performed on a Waters e2695 system (Waters, USA) equipped with a hypersil ODS column (Elite, Dalian, China). The mobile phase A was a mix of natrium aceticum (pH 4.8), water and tetrahydrofuran with the ratio of 410:85:5, and mobile phase B was pure acetonitrile. Flow rate was 1ml/min. The column temperature was set at 30°C. Samples were detected at the wavelength of 265nm. Injection volume was 20μL.

Immunofluorescent staining

After intracardiac perfusion with normal saline followed by fixation with 4% cold paraformaldehyde (PFA), the rat brains were separated and postfixed in 4% PFA for 6 h at 4 °C. Then the brains were incubated in 30% sucrose- 100 mM sodium phosphate buffer (pH 7.4) for 48h at 4 °C. Serial coronal sections (20 μm) were made from the bregma anterior-posterior −2.0 mm to −3.30mm. Totally 60 sections were collected per brain. At least three sections per brain were selected for NKB, Kisspeptin, GnRH/NK3R and GnRH/Kiss1r immunofluorescent staining, respectively.
For immunofluorescence, the primary antibodies rabbit anti-NKB (1:1000, Novus biologicals, NB300-201), rabbit anti-Kisspeptin (1:1000, H-048-56, Phoenix Pharmaceuticals), rabbit anti-Kiss-1r (1:500, AKR-001, Alomone Labs), rabbit anti-NK3r (1:500, abx217136, Abbexa) and mouse anti-GnRH (1:1000, MAB5456-C, Millipore, Billerica, MA) were used. After incubating with the antibodies for 24h at 4 °C, the sections were washed with PBS and then were treated with goat anti-mouse IgG (H + L) Alexa Fluor ®555 or 488 - conjugated or goat anti-rabbit IgG (H + L) Alexa Fluor ®488 (Invitrogen, Eugene, OR, USA) secondary antibodies. According to the manufacturer’s instructions, DAPI (Beyotime Biotechnology, Shanghai, China) was used to label nucleus. For negative controls, sections were incubated with PBS instead of the primary antibodies. Fluorescence images were captured using a Zeiss Axioskop 40 microscope(Carl Zeiss, Oberkochen, Germany). Image Pro-Plus 6.0 software was used for semi-quantitative measurement of immunofluorescent density. Values (three slides for each brain) of integral optical density (IOD) in individual cells represented the quantity of objective protein and were calculated using the following equation: $\sum \text{IOD}/\sum \text{DAPI}$.

**Statistical analysis**

All statistical analyses were performed with SPSS software (version 16.0), and the data were analyzed using the one-way Analysis of Variance (ANOVA). The data were expressed as the mean ± s.e.m. Statistical significance was set at $P<0.05$ for all tests.

**Results**

**Troxerutin reduced body weight, improved abnormal ovarian morphology and function in PCOS rats**

We first investigated whether troxerutin administration has any influence on body weight in DHT-induced PCOS rat models. The time course of troxerutin administration and evaluation of phenotypes were illustrated (Figure 1). Body weight for individual animal was weighed weekly after DHT implantation for up to 4 weeks. Body weight at baseline and 1st week showed no significant difference among all groups. DHT-treated rats showed significantly higher body weight at 2nd ($P<0.01$), 3rd ($P<0.01$) and 4th ($P<0.001$) week than the age-matched sham rats which denoted normal body weight, indicating that PCOS group have gained more body weight, hence a PCOS-like over weight feature. In comparison to PCOS group, treatment with troxerutin 150mg/kg did not induce any significant changes. Nevertheless, the body weight of PCOS+300mg/kg troxerutin group were markedly reduced as compared to that of PCOS group at 2nd ($P<0.05$), 3rd ($P<0.01$) and 4th ($P<0.05$) week, suggesting that troxerutin alleviated increased body weight in PCOS. Since ovarian histological change is another feature of PCOS, we also investigated whether troxerutin improved histological structure in rat ovaries. As demonstrated in figure 1b and 1c, normal ovarian histological features were observed in sham sections, which generally exhibited corpora lutea and follicles in various stages of development. In ovarian sections of PCOS rats, microscopic examination confirmed the absence of corpora lutea and the presence of cystic follicles ,which were characterized by a large fluid-filled structure with an attenuated granulosa cell layer and a thickened theca internal cell layer. We furthered calculated the percentage of follicles of total and the results
showed that PCOS rats exhibited significantly increased primordial and decreased primary follicles as compared to sham rats ($P<0.001$; $P<0.001$), and troxerutin 300mg/kg inhibited the markedly low percentage of primary follicles in PCOS rats ($P<0.05$). As compared to the multi-cystic follicles in ovaries of PCOS rats, the number of cystic follicles in troxerutin groups showed tendency toward decreasing. As figure 1.d demonstrated, sham group showed normal and stable estrous cycle, DHT induced cycle disruption in PCOS group, troxerutin administration 300mg/kg in PCOS rat models prevented and substantially reversed the impaired estrous cycle. These findings suggest that troxerutin is a positive regulator of increased body weight, abnormal ovarian histological structure and ovary function in PCOS rats.

**Troxerutin reversed the abnormal serum levels of gonadotrophin and testosterone in PCOS rats**

Serum LH, FSH and testosterone levels were measured at the fourth week by ELISA (Figure 2). PCOS rats showed a significant increase in serum LH and testosterone levels compared to sham rats ($P<0.001$; $P<0.05$), which was in accordance with the characteristic LH and testosterone elevation in patients with PCOS. Though there was no significant difference in FSH levels among all four groups, administration of troxerutin to rats for 4 weeks significantly reversed the elevated serum LH (300mg/kg; $P<0.01$) and testosterone levels (both 150mg/kg and 300mg/kg; $P<0.05$), suggesting that troxerutin has a beneficial effect on the aberrant gonadotrophin and testosterone in PCOS rats.

**Troxerutin inhibited the elevated GnRH levels in PCOS rats**

The origin of LH and FSH alteration often times lies at hypothalamic GnRH level, which plays a pivotal role in stimulating pituitary release of gonadotrophin, thus we further detected the serum and hypothalamic-pituitary status of GnRH. As shown in figure 3, the western blotting results showed increased GnRH in hypothalamic-pituitary of the PCOS rats ($P<0.01$), however, troxerutin 300mg/kg markedly decreased the level of GnRH compared to the PCOS rats ($P<0.05$; Figure 3b).

**Troxerutin altered GnRH regulatory neurotransmitters in the hypothalamus of PCOS rats.**

GnRH could be influenced by regulatory neurotransmitters such as the major inhibitory GABA and stimulatory Glu. As depicted in figure 4, neurotransmitter levels in the hypothalamus showed significant difference among certain groups. To be specific, PCOS rats exhibited markedly low GABA ($P<0.01$) and high Glu ($P<0.01$) compared to sham rats. Although no significant difference was observed between troxerutin 150kg/mg group and PCOS group, administration of troxerutin 300kg/mg to rats for 4 weeks successfully reversed the notably low GABA($P<0.05$) and high Glu ($P<0.01$) levels in the hypothalamus compared to PCOS group.

**Troxerutin reduced the expression of Kisspepetin1/Kiss1r and Neurokinin B/NK3r in the hypothalamus**

Kisspeptin and NKB secreted from the KNDy neurons in ARC, are considered to be novel GnRH stimulatory neurotransmitters and stimulate expression of kisspeptin receptor (kiss1r) and neurokinin B receptor (NK3r) of GnRH neurons in the median eminence (ME). Here, we carried out immunofluorescence
staining to observe whether troxerutin administration alters the expression of kisspeptin/kiss1r and NKB/NK3r in their respective regions (Figure 5 and Figure 6). The IOD of positive cells in the ARC were markedly increased in PCOS rats compared to sham rats (P<0.001; P<0.001), troxerutin 300mg/kg treatment significantly reversed the increase of kisspeptin and NKB IOD induced by DHT (P<0.05; P<0.01). Meanwhile, the changes of the expression of kiss1r and NK3r in ME were also observed. The numbers of GnRH/kissr1/NK3r positive cells increased in the PCOS group compared to the sham group(P<0.001; P<0.01; P<0.01), however, they were significantly decreased in the 300mg/kg troxerutin-treated rat (P<0.01; P<0.05; P<0.05) respectively.

Discussion

PCOS is the most common endocrine condition affecting approximately 20% of reproductive-aged women. Though some ovulation induction agents, such as letrozole, clomiphene citrate and metformin, have been used to improve reproductive outcomes in clinic, the side effects of these drugs should not be ignored[24, 25]. Thus, numerous studies attempted to guide the development of new and effective therapies for PCOS. Recently, treatment based on traditional Chinese medicine and food ingredients, such as AF extract, soy isoavones and crocetin, provided a novel therapeutic way for PCOS [26-28]. All these observations suggested rooms for improvement in PCOS therapies. Possession of beneficial effects on animal models of different central nervous system diseases made troxerutin, a rutin derivative, an attractive therapeutic method for us to investigate its possible effect on PCOS. Specifically, troxerutin inhibited cyclin-dependent kinase 1 expression, enhanced type 1 protein phosphatase α dephosphorylation and abolished MEK/ERK1/2/C/EBP β activation, which subsequently reversed the memory impairment in the Domoic acid-treated mice[20]. The neuroprotective potential of troxerutin in 6-OHDA rat model of Parkinson's disease was through mitigation of apoptosis, astrogliosis, oxidative stress and part of its effect was dependent on PI3K/ERβ signaling[21]. Troxerutin and cerebroprotein hydrolysate injection acted as a neuroprotective agent against cerebral ischemia/reperfusion injury via anti-inflammation, anti-apoptosis and blood-brain barrier maintenance[29]. In the present study, we evaluated the possible effect of troxerutin for up to four weeks in a DHT-induced PCOS rat model, and two important findings were revealed. First, administering troxerutin 300mg/kg exerted a beneficial effect of reducing body weight, the elevated levels of LH, testosterone and GnRH in PCOS rats and its effect was superior to troxerutin 150mg/kg. Second, the potential mechanism behind the observed effect on increased body weight, polycystic ovaries, impaired estrous cycle and endocrine aberration involved altering GABA, glutamate, kisspeptin/kiss1r and NKB/NK3r in the hypothalamic pituitary region. To the best of our knowledge, this is the first study to investigate the possible effect of troxerutin on PCOS using a rat model and the potential mechanism of its protective effect may be via regulating GABA, glutamate, kisspeptin/kiss1r and NKB/NK3r in the brain/pituitary.

GnRH pulses stimulate the synthesis and secretion of LH and FSH from anterior pituitary. It is well known that although produced in the same cell named gonadotroph, LH and FSH synthesis is regulated by different frequency of GnRH pulses, with LH favored by fast pulse frequencies(>1 pulse per hour) and FSH favored by slow pulse frequencies(<1 pulse per 2-3 hour). As for PCOS, a neuroendocrine hallmark is
persistent and rapid GnRH pulses, which favor pituitary synthesis of LH and contribute to the increased LH levels[30]. LH is a stimulus for androgen synthesis, so the increased LH drive in turn caused the elevated androgens [31, 32]. Therefore, in terms of biochemical indicators, the most important and characteristic abnormality referring to PCOS is elevated LH, subsequently elevated testosterone and low to normal FSH levels in serum[4, 33]. Consistent with previous studies, PCOS rats displayed similar biochemical abnormality including markedly increased LH, concomitantly increased testosterone levels and non-significant FSH levels as compared to sham rats. Next, we observed the serum LH, testosterone and FSH levels in PCOS and troxerutin groups in order to determine whether troxerutin treatment has any therapeutic effect on PCOS. As the results revealed, troxerutin treatment indeed inhibited the elevated LH and testosterone levels without significant influence of FSH level. In addition, we further estimated the hypothalamic status of GnRH, which was significantly increased in PCOS rats. However, there was a significant troxerutin-caused (300mg/kg) decline in hypothalamic GnRH. In regard to the serum hormone levels in rats, many previous studies have reported that LH were generally in the range of 1.5~3 mIU/ml[34, 35], FSH levels varied from 8mIU/ml to 20mIU/ml [26, 36] and testosterone varied from 0.2 to 10 ng/ml[34, 36-39], which were approximately consistent with the corresponding results in the present study. Conclusively, troxerutin treatment 300mg/kg to rats for up to 4 weeks inhibited the hyperactive GnRH/LH system in PCOS rats.

Compelling evidence provided the strong support for brain being the culprit in PCOS[40, 41]. The secretion of GnRH from the brain is itself regulated by numerous upstream factors, such regulators include GABA, norepinephrine, dopamine, serotonin and glutamate, amongst others[42, 43]. While specific impairments in the brain were difficult to assess in humans, rat models greatly facilitated the identification of difference underpinning the hyperactive HPG axis in PCOS. Therefore, the representative and GnRH regulators of inhibitory GABA and stimulatory Glu were observed in the present study. In comparison to sham rats, PCOS rats showed a turnover of regulatory neurotransmitters with a significantly decreased GABA and increased Glu, which were both reversed by administering troxerutin 300mg/kg to rats for up to 4 weeks. As far as the hypothalamic neurotransmitter levels were concerned, previous studies reported that hypothalamic GABA was in range of 12-20μM/g and hypothalamic Glu was in range of 22-45μM/g in rats, each approximately consistent with our findings after unit conversion. To date, one arcuate nucleus population of particular recent interest has been the kisspeptin/neurokinin B/dynorphin expressing 'KNDy' neurons, which synthesized and released GnRH stimulatory neurotransmitters kisspeptin and NKB. Protrusions of GnRH neurons in the ME express their receptors kiss1r and NK3r. In the present study, troxerutin decreased the expression of kisspeptin1 and NKB in the arcuate nucleus, and also the receptors kiss1r and NK3r in GnRH positive neurons of the ME.

Declarations

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Authors' Contributions

Zixuan Gao and Xiaochen Ma performed the experiments, collected and analyzed the data and prepared the manuscript. Yuhang Ge, Lei Wang and Ping Fu performed the experiments and provided the data. Zhian Liu, Ruiqin Yao, Xiaonan Yan designed the study, provided financial support and edited the manuscript. All the authors approved the final approval of the manuscript.

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Availability of data and materials

All the data is contained in the manuscript.

Ethics approval

All animal experiments were approved by the Institutional Animal Care Committee of Xuzhou Medical University and in strict accordance with the Guidelines for the Care and Use of Laboratory Animals(National Research Council of People's Republic of China,2010)

Consent for publication

Not applicable.

Competing interests

There is no conflict of interest to declare. The funders had no role in the study design; in the collection, analyses or interpretation of data; in the manuscript and decision to publish the results.

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Conclusions
Taken above, these results indicated that the protection of troxerutin against PCOS partially might be due to its ability to regulate hypothalamic GABA, Glu, kisspeptin/kiss1r and NKB/NK3r. Although it remains determined whether this troxerutin-caused reversal is permanent, the present study highlighted its therapeutic potential for PCOS. More in depth research will be necessary to determine whether troxerutin targeted at hypothalamic neurotransmitters alone is a promising therapeutic approach for the treatment of PCOS.

**Abbreviations**

ANOVA: Analysis of Variance; ARC: arcuate nucleus; DHT: Dihydrotestosterone; ELISA: Enzyme Linked Immuno Sorbent Assay; FSH: follicle stimulating hormone; GABA: Gamma-aminobutyric acid; Glu: glutamate; GnRH: Gonadotrophin-releasing hormone; H&E: hematoxylin and eosin; HPLC: High-performance liquid chromatography; HPG: hypothalamus-pituitary-gonad; IGF-1: Insulin growth factor-1; IOD: integral optical density; Kiss1r: Kisspeptin 1 receptor; KNDy: kisspeptin-/ neurokinin B-/ dynorphin; LH: luteinizing hormone; ME: median eminence; NGF: nerve growth factor; NKB: neurkinin B; PCOS: Polycystic ovary syndrome; PFA: paraformaldehyde.

**Declarations**

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**Availability of data and materials**

The datasets used and/or analyzed in the present study are available from the corresponding author on reasonable request.

**Authors’ contributions**

RQ Y and XN Y conceived and designed the idea to this paper; ZX G, XC M, YH G, L W, P F, J L collected and analyzed the data, and drafted the paper; ZA L, RQ Y and XN Y analyzed the data and revised the final paper. All authors read and approved the final manuscript.

**Ethics approval and consent to participate**
The experiments were conducted in accordance with the Regulation of Animal Research Ethics in China. The ethical approval was granted by the Xuzhou Medical University Animal Research Ethics Committee (Project number 201811W011, Xuzhou, China).

**Consent for publication**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

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**Figures**

**Figure 1**

Troxerutin reduced obesity and improved abnormal ovarian morphology in PCOS rats (a) Body weight (b) Hematoxylin and eosin staining of representative ovaries. For a, P values were determined by one-way ANOVA with Turkey’s multiple comparison test and data were presented as means±s.e.m. n=12 rats per group. *P<0.05, **P<0.01, ***P<0.001 vs. sham group; #P<0.05, ##P<0.01 vs. PCOS group. For b, follicles in various stages of development were indicated by black triangles, corpora lutea were indicated by asterisks, while the cystic follicles were indicated by hashtags. Scale bar: 200μm. Images were representative of three independent experiments with similar results.
Figure 2

Troserutin reversed the abnormal serum levels of gonadotrophin and testosterone in PCOS rats (a) Serum LH levels (b) Serum FSH levels (c) Serum testosterone levels. P values were determined by one-way ANOVA with Turkey’s multiple comparison test and data were presented as means±s.e.m. n=9 rats per group. *P<0.05, **P<0.001 vs.sham group; #P<0.05, ##P<0.01 vs. PCOS group.

Figure 3
Troxeutin reduced the elevated GnRH levels in PCOS rats (a) Serum levels of GnRH (b) Hypothalamic status of GnRH. P values were determined by one-way ANOVA with Turkey’s multiple comparison test and data were presented as means±s.e.m. n=9 rats per group.*P<0.05, **P<0.01, ***P<0.001 vs. sham group; #P<0.05, ##P<0.01 vs. PCOS group; △△P<0.01 vs. PCOS+150mg/kg troxerutin group.

![Graphs showing changes in GnRH levels and neurotransmitters](image)

**Figure 4**

Troxeutin altered GnRH regulatory neurotransmitters in the hypothalamus of PCOS rats (a) Hypothalamic levels of GnRH inhibitory GABA (b) Hypothalamic levels of GnRH stimulatory glutamate. P values were determined by one-way ANOVA with Turkey’s multiple comparison test and data were presented as means±s.e.m. n=9 rats per group. **P<0.01 vs. sham group; #P<0.05, ##P<0.01 vs. PCOS group.
Figure 5

Troserutin reduced the expression of Kisspeptin1/Kiss1r and Neurokinin B/NK3r in the hypothalamus (a) and (b) Immunofluorescence photomicrograph showing that the expression of Kisspeptin1 and NKB in the arcuate nucleus (ARC) and Kiss1r, NK3r in GnRH positive neurons in the median eminence (ME). Scale bar, 50 μm. (c) and (d) Quantitative analysis integral optical density (IOD) of Kisspeptin1 and NKB in the ARC and the number of Kiss1r/ GnRH and NK3r/GnRH positive cells in the ME. P values were determined
by one-way ANOVA with Turkey's multiple comparison test and data were presented as means±s.e.m. n=6 rats per group. **P<0.01, ***P<0.001 vs. sham group; #P<0.05, ##P<0.01 vs. PCOS group.