Evaluating the Therapeutic Efficacy of Si-Wu-Tang Decoction and Concentrated Extract in Follicular Maldevelopment-Related Menstrual Disorders Through Pharmacokinetic/Pharmacodynamic Studies

Chia-Jung Lee1,2,3, Alinafe Magret Kapelemera2, Yi-Zhe Tsai1, Ching-Tzu Lee4, Ming-Yi Xu5 and Ching-Chiung Wang1,2,3,6*

1 Graduate Institute of Pharmacognosy, College of Pharmacy, Taipei Medical University, Taipei, Taiwan, 2 PhD Program in Clinical Drug Development of Herbal Medicine, College of Pharmacy, Taipei Medical University, Taipei, Taiwan, 3 Traditional Herbal Medicine Research Center, Taipei Medical University Hospital, Taipei, Taiwan, 4 Department of Traditional Chinese Medicine, Taipei Municipal Wanfang Hospital, Taipei, Taiwan, 5 School of Medicine, College of Medicine, Taipei Medical University, Taipei, Taiwan, 6 School of Pharmacy, College of Pharmacy, Taipei Medical University, Taipei, Taiwan

Si-Wu-Tang (SWT), a traditional Chinese formula, is commonly used to relieve menstrual discomfort and climacteric syndrome. Water decoction (WD) and concentrated herbal extract (CHE) are the two most common formulations of traditional Chinese medicine (TCM). However, few studies have reported the equivalency of these two formulations. In this study, 23 healthy volunteers were included to determine the pharmacokinetic (PK) equivalent dosage of WD and CHE, and 25 infertile women with follicular maldevelopment to evaluate the pharmacodynamic (PD) effects on menstrual disorders. The randomized, two-way crossover comparative PK study of SWT-WD and SWT-CHE analyzed the active component, ferulic acid. The results showed that clinical doses of 170 mL SWT-WD and 18 g SWT-CHE produced the same amount of ferulic acid in the blood. The PD study showed that patients who took both of these formulations had an initial luteinizing hormone/follicle-stimulating hormone ratio of <1; however, the value returned to normal and their symptoms all improved after taken SWT. Our results showed that WD and CHE, both prepared from 40 g of SWT, displayed bioequivalence upon PK/PD analysis.

Keywords: Si-Wu-Tang, ferulic acid, menstrual disorders, follicular maldevelopment, pharmacokinetic/pharmacodynamic

INTRODUCTION

Traditional Chinese medicine (TCM) has been developed and used for over 2,500 years. TCM is still widely used in Taiwan and is beginning to receive global recognition. A decoction is an important dosage form of TCM. Boiling has a great influence on its clinical efficacy. Decoctions are difficult to prepare and transport, and it is challenging to maintain quality control. Moreover, a commonly commercial
According to TCM theory, SWT is a foundational blood tonic (Ministry of Health and Welfare, 2018). In addition, Japan also has Pharmacy and Taiwan Herbal Pharmacopeia, recommended (August 31, 1995) and No. 89037929 (June 29, 2000) from Ministry consistent quality. According to the announcement No. 84056272 water, temperature, and time taken to produce products with extracts (CHEs) have been used since 1963. Depending on the advancement of pharmaceutical technology, concentrated herbal resources and may reduce clinical ef.

Ingredients (Schulzki et al., 2017). This results in loss of medicinal ingredients (Lai et al., 2013). Nevertheless, many studies have not systematically explained the specific effects of SWT on blood and menstruation. In the case of two different pathological phenomena, how the herbs exert their synergistic or additive effects, with one side having multiple effects is unexplained.

In this study, we cooperated with Taiwan’s good manufacturing practice (GMP) that drug manufacturers use to produce CHEs and used the same source of medicinal materials to prepare a decoction of SWT. To explore the bioequivalence of the two formulations, we used healthy volunteers and performed pharmacokinetic (PK) and pharmacodynamic (PD) studies to observe follicular development. Subjects took both preparations in order to evaluate the bioequivalence of the different formulations. We hope that the results will provide a reference for future clinical use.
MATERIAL AND METHODS

Preparation and Quality Control of SWT
Preparation of SWT-WD and SWT-CHE

The SWT prescription was based on the unified formula announced by the Committee on Chinese Medicine and Pharmacy of the Department of Health (Taipei, Taiwan) (Ministry of Health and Welfare, 2018). Angelica Radix (Angelica sinensis (Oliv.) Diels, Dang Gui), Chuanxiong Rhizoma (Ligusticum chuanxiong Hort., Chuan Xiong), Paeoniae Radix (Paeonia lactiflora Pall., Shao Yao), and Rehmanniae Radix (Rehmannia glutinosa Libosch., Shu Di Huang) were purchased from Sun Ten Pharmaceutical (New Taipei City, Taiwan). The medicinal materials were authenticated by the non-profit organization, Brion Research Institute of Taiwan. Voucher specimens (No. AS-20180001 for Angelica Radix, No. LC-20180001 for Chuanxiong Rhizoma, No. PL-20180001 for Paeoniae Radix and No. RG-20180001 for Rehmanniae Radix) was deposited at the College of Pharmacy, Taipei Medical University. The SWT prescription included the following herbs: 600 g of each herb was weighed (total drug weight of 2,400 g), placed in an extractor, and 20 L of filtered water was added and boiled for 30 min until half the volume was left (Tseng et al., 2006). Ten liters of SWT-WD was packaged into 60 packets and the volume of each bag was 170 mL. One packet was used each morning and evening (twice a day). SWT-CHE was purchased from Sun Ten Pharmaceutical (New Taipei City, Taiwan). The batch number of SWT-CHE was 140243. Each 15 g of SWT-CHE contained 8.5 g of extract and 6.5 g of cornstarch conforming to the drug permit license. An 8.5 g extract was derived from the four herbs at a ratio of 1:1:1:1 and each drug was 7.5 g (total drug weight present as 30 g). Based on Sun Ten Pharmaceutical prescription instructions, 30 g of dry herbs was made into 15 g SWT-CHE granules and 18 g of granules was the recommended dose per day. As for the conversion, based on the daily dose of the SWT-CHE we calculated the amount of herbal materials to be 36 g/day. Therefore, the SWT-WD used 170 mL to perform the following experiment, which was the equivalent of 40 g of herbal materials.

High-Performance Liquid Chromatography (HPLC)
Analysis of Marker Substances in SWT-WD and SWT-CHE

The HPLC apparatus was composed of an SCL-10Avp System Controller, an LC-10ATvp Liquid Chromatograph Pump, an SPD-M10A Diode Array Detector, an SIL-10AVp Auto Injector, a CTO-10A Column Oven, FCV-10Avp Flow-Channel Selection Valves (all from Shimadzu, Tokyo, Japan), and an ERC-3415 Degasser (ERC, Altegolfsheim, Regensburg, Germany). The stationary phase consisted of a Purospher® STAR RP-18e reversed-phase column (5 μm, 4 mm i.d. x 250 mm, Merck, Germany). The mobile phase for chromatographic separation of (A) was 0.05% (v/v) formic acid aqueous solution and (B) acetonitrile using a gradient elution of 12% B at 0–15 min, 17% B at 15.01–30 min, and 100% B at 30.01–60 min. The flow rate was 1.0 mL/min, and the column temperature was maintained at 40°C. We used an ultraviolet wavelength of 320 nm to detect the contents of ferulic acid in SWT-WD and SWT-CHE. An aliquot of 1 g of SWT-CHE powder from the above pharmaceutical company was extracted ultrasonically with 10 mL of water for 1 h at room temperature. The extracted solution was centrifuged at 13,000 rpm for 10 min. The supernatant was collected and diluted 50-fold. Further, SWT-WD was diluted 50-fold with distilled water. The sample solutions (SWT-WD and SWT-CHE) were then filtered through a 0.45 μm filter and 10 μL was directly injected into the HPLC system. Calibration curves were plotted based on the analyte peak areas: $y = 11762x - 4380.7$, and $R^2 = 0.9938$.

PK Study of SWT

Subjects

All experimental protocols were reviewed and approved by the Institutional Review Board (IRB) of Taipei Medical University, Taiwan (no.: 100024). This study recruited 23 non-menstrual healthy women (aged 20–35 years). The inclusion criterion was that volunteers could take no nutritional supplements (such as vitamins or iron agents) or drugs (including Chinese and Western medicine) for one week before the experiment.

Study Design

The study had an open-label, two group crossover treatment design. Twenty-three volunteers were separated into two groups. In Group A, 14 volunteers were given 18 g of SWT-CHE and 135 mL of SWT-WD. In Group B, 9 volunteers were given 18 g of SWT-CHE and 170 mL of SWT-WD. Volunteers could choose to take SWT-CHE or SWT-WD. A 28-day washout period was observed between doses. For each group, volunteers returned to the clinical facility for subsequent blood drawing for PK sampling.

PK Sampling

In each group, blood samples for determination of ferulic acid were collected during confinement at the clinic prior to drug administration and at 0, 10, 30, 60, 120, and 180 min after administration. Five milliliter of blood were collected for each time point from median cubital vein. All blood samples were drawn into blood collection tubes, the interior of which was coated with dipotassium ethylenediaminetetraacetic acid (EDTA).

Analytical Methods

Blood samples were cooled in an ice bath and separated by centrifugation at 2000 × g for at least 10 min at 4°C. Two aliquots of a minimum of 1 mL plasma were dispensed into polypropylene tubes and maintained at −20 ± 5°C. Plasma samples were used to determine ferulic acid concentrations in human plasma with EDTA as an anticoagulant using an HPLC.
PK Analysis
PK analysis was performed using Phoenix WinNonlin ver. 6.4 (Certara). The maximal measured plasma concentration (C_{max}) and the time at which the maximal plasma concentration was observed (T_{max}) were taken from the observed plasma concentration-time profile. The area under the concentration-time curve from time zero until the last measurable concentration or the last sampling time, t, whichever occurred first (AUC0-t), was calculated using the trapezoidal method.

PD Study of SWT in Humans
According to the PK equivalence of the ferulic acid experiment in the first stage, we found that 170 mL of SWT-WD was equivalent to 18 g of SWT-CHE. Therefore, two doses were provided for further PD studies. Twenty-five volunteers were diagnosed at Wan-Fang Hospital, using an ultrasonic examination, with an abnormal LH/FSH ratio, abnormal menstrual cycles, and follicular maldevelopment. Together, based on TCM, Dr. Ching-Tzu Lee diagnosed patients who were treated with SWT and who were included in this project. Before taking SWT, blood was taken to check the estradiol (E2), FSH, LH, and progesterone (P4), testosterone (T) levels, and an ultrasonic examination was performed. Volunteers were randomly selected to take 170 mL of the SWT-WD, which was equivalent to 18 g of SWT-CHE, according to their own wishes. A parallel design was used. During the experiment, blood was collected each month (if the menstrual period occurred, blood was collected before the 3rd–5th day), and the observation point was at 3 months. At the end of the experiment, follicular development was detected using ultrasound. During the experiment, subjects were not allowed to take other SWT-related products, iron-containing drugs, or other TCM products.

RESULTS
Ferulic Acid Contents in SWT-WD and SWT-CHE Groups
According to the above data, the same concentration of ferulic acid in SWT-WD and SWT-CHE were calculated as previously described in the Methods section. Each gram of SWT-CHE contained 58.5 mg of ferulic acid, and each milliliter of SWT-WD contained 440 µg of ferulic acid. After calculation, 18 g of SWT-CHE and 135 mL of SWT-WD were determined to have the same amount of ferulic acid (Figure 1).

PK Study of SWT
The 23 volunteers' plasma ferulic acid concentration-time courses following a single administration are shown in Figure 2. Group A was given 18 g of SWT-CHE and 135 mL of SWT-WD that presented the same amount of ferulic acid. Group B was administered 18 g of SWT-CHE and 170 mL of SWT-WD that presented the clinical dosage. These two groups were used to compare the bioequivalence of ferulic acid. The derived values of the PK parameters are presented in Table 1. Results showed that the maximum concentration (C_{max}) and the area under the curve (AUC_{0-180} min) of 18 g of SWT-CHE were higher than those with 135 mL of SWT-WD with the same amount of ferulic acid compared to group A. Further, in the clinical dosage group B, C_{max} and the AUC_{0-180min} presented similar data without a significant difference. There was no significant difference between CHE and WD at T_{max} and T_{1/2}. Therefore, we chose Group B, 170 mL of SWT-WD and 18 g of SWT-CHE, for using as the clinical dosage for the ovarian follicular maldevelopment PD bioequivalence study.

PD Study of SWT
This study was hosted in consultation with Chinese and Western medicinal specialists at Wan-Fang Hospital. Patient inclusion criteria according to clinical conditions were as follows: (1) an LH/FSH ratio of >1, (2) abnormal LH or FSH levels, and (3) follicles <8 mm and their number greater than 12; also a patient mentioning menstrual abnormalities (cyclic disorder, or abnormal menstrual blood color or quantity). Twenty-five patients were included in this study; 12 patients underwent SWT-WD and 13 patients underwent SWT-CHE. However, 12 patients withdrew from the process for no given reason, one patient withdrew due to diarrhea (an expected side effect), and
FIGURE 1 | HPLC chromatogram of ferulic acid in (A) Si-Wu-Tang (SWT)-water decoction (WD) and (B) SWT-concentrated herbal extract (CHE).

FIGURE 2 | Area under the curve (AUC)-time curve of ferulic acid in the subject’s serum. (A) Group with 135 mL of the Si-Wu-Tang (SWT)-water decoction (WD) and 18 g of the SWT-concentrated herbal extract (CHE) and (B) the group with 170 mL of SWT-WD and 18 g of SWT-CHE.
one patient changed to another Chinese medicine. Ultimately, six patients completed SWT-WD and five patients completed SWT-CHE.

Before taking SWT, endocrine hormones were determined in all patients with ovarian follicular maldevelopment; FSH and LH levels were found to be significantly higher, and E₂ levels were found to be significantly lower; this is considered normal levels for the menopausal period. Due to the low level of estrogen, the level of emergence of bottom cells was also low.

After taking SWT-WD or SWT-CHE, the LH of patients was significantly lower in the first month. As the duration of taking SWT increased, the effect was more significant (Figure 3A) and there was no significant difference between SWT-WD and SWT-CHE. In the estradiol analysis, SWT-WD reduced estradiol levels by the third month and SWT-CHE reduced estradiol levels by the second month (Figure 3C). However, the FSH and progesterone levels of patients exhibited no significant difference before and after taking SWT-WD or SWT-CHE (Figures 3B, D). Taking SWT-WD reduced testosterone levels in the first and second months. The LH/FSH ratio of all six patients was <1 and the value returned to normal after the study (Figure 3F). In the SWT-CHE group, the LH/FSH ratio of four

### TABLE 1 | Pharmacokinetic parameters of ferulic acid in serum after taking Si-Wu-Tang.

| Formulation | Dosage | T<sub>max</sub> (min) | C<sub>max</sub> (ng/mL) | AUC<sub>0-180 min</sub> (min × ng/mL) | Cl (mL/min) | T<sub>1/2</sub> (min) |
|-------------|--------|-----------------------|----------------------|---------------------------------------|-------------|------------------|
| A           | 135 mL | 25.7 ± 8.5<sup>a</sup> | 30.9 ± 8.1<sup>a</sup> | 3619.7 ± 965.6<sup>a</sup> | 2310.4 ± 531.1<sup>a</sup> | 131.7 ± 74.5<sup>a</sup> |
|             | 18 g   | 32.9 ± 12.7<sup>a</sup> | 42.1 ± 14.2<sup>b</sup> | 4404.7 ± 1030.5<sup>b</sup> | 1892.7 ± 474.8<sup>b</sup> | 88.2 ± 46.7<sup>b</sup> |
| B           | 170 mL | 31.1 ± 12.7<sup>a</sup> | 46.6 ± 25.1<sup>b</sup> | 4276.2 ± 1140.0<sup>b</sup> | 2463.4 ± 642.5<sup>b</sup> | 156.9 ± 79.7<sup>b</sup> |
|             | 18 g   | 30.0 ± 0<sup>a</sup>   | 44.0 ± 11.6<sup>a</sup> | 4046.1 ± 752.5<sup>a</sup>  | 2019.5 ± 409.7<sup>a</sup> | 78.4 ± 12.7<sup>a</sup> |

<sup>a</sup>Time at which C<sub>max</sub> was observed; <sup>b</sup>maximal measured plasma concentration; AUC, area under the concentration-time curve; Cl, clearance; T<sub>1/2</sub>, half-life.

**FIGURE 3** (A) Luteinizing hormone (LH), (B) follicle-stimulating hormone (FSH), (C) estradiol, (D) progesterone, (E) testosterone, and (F) the LH/FSH ratio in the serum of subjects in the Si-Wu-Tang (SWT)-water decoction (WD) and SWT-concentrated herbal extract (CHE) treatment groups.
DISCUSSION

Ferulic acid, a quality component of *A. sinensis*, and *L. chuangxiong*, has also been used as a quality component to evaluate the amount in SWT-WD and SWT-CHE. Based on the analysis of data from group A, 135 mL of SWT-WD and 18 g of SWT-CHE represented the same amount of ferulic acid, and according to Sun Ten Pharmaceutical prescription instructions, 170 mL of SWT-WD and 18 g of SWT-CHE represented the clinical dosage. However, when 18 g of SWT-CHE was taken, the AUC of ferulic acid was higher than that with 135 mL of SWT-WD, but was similar to that with 170 mL of SWT-WD. In the experiment with equal amounts of ferulic acid, SWT-CHE presented a higher C\text{max} and AUC than SWT-WD. This showed the superiority of the concentrated extract with a high blood concentration. According to Figure 2 and Table 1, FA was shown quickly absorbed after oral administration and reaches the peak plasma concentration (C\text{max}) around 30 min (T\text{max}). Ferulic acid was metabolism in the liver through CYP 1A1 and 2B7 of the UDP-glucuronosyl-transferase (UGT) (Li, X., Shang, L., Wu, Y., Abbas, S., Li, D., Netter, P., Ouzzine, M., Wang, H., Magdalou, J., 2011. Identification of the human UDP-glucuronosyltransferase isoforms involved in the glucuronidation). As clinical studies, glucuronide (3–20%) and sulfo-glucuronide (60–90%) are the most abundant metabolites of FA in plasma, whereas only a low percentage of unmodified FA (9–20%) has been found. FA and its metabolites are excreted mainly from the kidney. (Zhao et al., 2003; Zhao et al., 2004). Our results showed that both SWT-CHE and SWT-WD presented similar characteristic.

Chinese medicine treatment for infertility is divided into five major types of diseases: kidney yang-deficiency, kidney yin-deficiency, liver qi stagnation, blood stasis, and phlegm, and dampness types (Hsu et al., 2006). According to the results, both formulations of SWT could ameliorate follicular maldevelopment and irregular menstrual cycles. It was speculated that patients with abnormal LH and FSH levels may have liver qi stagnation. When adjusting the daily dose of SWT for 3 months, either taking SWT-WD or SWT-CHE, both formulations produced sufficient blood concentrations of ferulic acid, which could improve ovarian follicular maldevelopment. It has been reported that oxidation could be a reason for causing failure of ovary function (Wang et al., 2018). Therefore, both agents presented equivalent effects. Taken together, ferulic acid is a good indicator to analyze the bioequivalence in both PK and PD studies. Despite the preclinical evidence supporting the potentially important role of ferulic acid in free radical-induced diseases. Whereas only a few clinical studies were performed keeping in mind a pharmacodynamic target or a specific disease. Our experiment showed that both SWT-CHE and SWT-WD with same amount ferulic acid could improve ovarian follicular maldevelopment.

However, from current clinical experience, doctors usually advise patients to use a decoction once in the morning and evening. According to our experiment, taking SWT-WD once a day, has already presented the same blood concentration as SWT-CHE and showed good amelioration of follicular maldevelopment. Therefore, the common daily dose of the decoction was twice that of the concentrated extract. Patients that have previously thought that the decoction was better than the concentrated extract may just have taken more drug than the concentrated extract. Our results showed that WD and CHE, both prepared from 40 g of SWT, displayed bioequivalence upon PK/PD analysis. Based on the results of this study, we recommend that the clinical dosage used in these two formulations should be considered as an equivalent amount between the PK and PD experiments (Figure 4). In conclusion, Si-Wu-Tang could improve ovarian follicular maldevelopment. But there are still many ingredients and related to the efficacy need to be extensively explored and discussed in future study.

![Figure 4](link)
DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation, to any qualified researcher.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Institutional Review Board (IRB) of Taipei Medical University, Taiwan (no.: 100024). The patients/participants provided their written informed consent to participate in this study.

REFERENCES

Chang, C. C., Lee, Y. C., Lin, C. C., Chang, C. H., Chiu, C. D., Chou, L. W., et al. (2016). Characteristics of traditional Chinese medicine usage in patients with stroke in Taiwan: a nationwide population-based study. J. Ethnopharmacol. 186, 311–321. doi: 10.1016/j.jep.2016.04.018

Cheng, C. Y., Kao, S. T., and Lee, Y. C. (2019). Ferulic acid ameliorates cerebral infarction by activating Akt/mTORC4/ERK-pBcl-2 anti-apoptotic signaling in the penumbra cortex following permanent cerebral ischemia in rats. Mol. Med. Rep. 19, 792–804. doi: 10.3892/mmr.2018.9737

Dai, Y., But, P. P., Chan, Y. P., Matsuoka, H., and Kubo, M. (2002). Antipruritic and antiinflammatory effects of aqueous extract from Si-Wu-Tang. Biot. Pharm. Bull. 25, 1175–1178. doi: 10.1248/bpb.25.1175

Fang, Z., Lu, B., Liu, M., Zhang, M., Yi, Z., Wen, C., et al. (2013). Evaluating the pharmacological mechanism of Chinese medicine Si-Wu-Tang through multi-level data integration. PLoS One 8, e72334. doi: 10.1371/journal.pone.0072334

Gao, J., Yu, H., Guo, W., Kong, Y., Gu, L., Li, Q., et al. (2018). The anticancer effects of ferulic acid is associated with induction of cell cycle arrest and autophagy in cervical cancer cells. Cancer Cell Int. 18, 102. doi: 10.1186/s12935-018-0595-y

Ghosh, S., Chowdhury, S., Sarkar, P., and Sil, P. C. (2018). Ameliorative role of ferulic acid against diabetes associated oxidative stress induced spleen damage. Food Chem. Toxicol. 118, 272–286. doi: 10.1016/j.fct.2018.05.029

Hanson, B., Johnstone, E., Dorais, J., Silver, B., Peterson, C. M., and Hotaling, J. (2017). Female infertility, infertility-associated diagnoses, and comorbidities: a review. J. Assist. Reprod. Genet. 34, 167–177. doi: 10.1007/s10815-016-0836-8

Hsu, H. Y., Hsu, C. S., and Hu, Q. F. (2006). Commonly Used Chinese Herb Formulas with Illustrations. 2nd edn (Irvine, California, USA: Oriental Healing Arts Institute).

Huang, G. C., Tsai, Y. Z., Lee, C. J., Chang, H. Y., and Wang, C. C. (2019). Observational studies on evaluating antioxidant capacity of the ovary in menopausal rats and in ovarian granulosa cell culture. Clin. Endocrinol. (Oxf.) 90, 579–585. doi: 10.1111/cen.13934

Lee, C. J., Hsueh, T. Y., Lin, L. C., and Tsai, T. H. (2014). Determination of protein-unbound rynchophylline branch distribution by microdialysis and ultra-performance liquid chromatography with tandem mass spectrometry. Biomed. Chromatogr. 28, 901–906. doi: 10.1002/bmc.3206

Lin, F. H., Lin, J. Y., Gupta, R. D., Toumas, J. A., Burch, J. A., Selim, M. A., et al. (2005). Ferulic acid stabilizes a solution of vitamins C and E and doubles its photoprotection of skin. J. Invest. Dermatol. 125, 826–832. doi: 10.1111/j.0022-202X.2005.23768.x

Lin, M. J., Chen, H. W., Liu, P. H., Cheng, W. J., Kuo, S. L., and Kao, M. C. (2019). The prescription patterns of traditional Chinese medicine for women with polycystic ovary syndrome in Taiwan: A nationwide population-based study. Med. (Baltimore) 98, e15890. doi: 10.1097/MD.0000000000015890

Ministry of Health and Welfare (2018). Taiwan Herbal Pharmacopoeia. 3rd edn. Committee. (Taipei, Taiwan).

AUTHOR CONTRIBUTIONS

C-JL and C-CW conceived and designed the experiments. C-JL, AK, Y-ZT, and C-TL performed the experiments. C-JL and M-YX analyzed the data. C-JL wrote the manuscript. C-CW revised the manuscript.

FUNDING

This work was financially supported by the by Taipei Medical University (Grant Number: TMU103-AE1-B38).

Okamoto, R., Sakamoto, S., and Noguchi, K. (1976). Effects of ferulic acid on FSH, LH and prolactin levels in serum and pituitary tissue of male rats. Nihon Naibunpi Gakkai Zasshi. 52, 953-958. doi: 10.1507/endocrine1927.52.953

Park, H. R., Kim, J. H., Lee, D., and Jo, H. G. (2019). Cangfu daotan decoction for polycystic ovary syndrome: a protocol of systematic review and meta-analysis. Med. (Baltimore) 98, e17321. doi: 10.1097/MD.0000000000017321

Ried, K. (2015). Chinese herbal medicine for female infertility: an updated meta-analysis. Complement Ther. Med. 23, 116–128. doi: 10.1016/j.ctim.2014.12.004

Schulzki, G., Nüßlein, B., and Sievers, H. (2017). Transition rates of selected metals determined in various types of teas (Camellia sinensis L. Kuntze) and herbal/ fruit infusions. Food Chem. 215, 22–30. doi: 10.1016/j.foodchem.2016.07.093

Sgarbona, A., Giacomazza, D., and di Carlo, M. (2015). Ferulic acid: a hope for Alzheimer’s disease therapy from plants. Nutrients 7, 5764–5782. doi: 10.3390/nu7070246

Sidra, S., Tariq, M. H., Farrukh, M. J., and Mobsin, M. (2019). Evaluation of clinical manifestations, health risks, and quality of life among women with polycystic ovary syndrome. PLoS One 14, e0223329. doi: 10.1371/journal.pone.0223329

Sun, J., Zhang, L., He, Y., Zhang, K., Wu, L., Fan, Y., et al. (2016). To unveil the molecular mechanisms of Qi and blood through systems biology-based investigation into Si-Jun-Zi-Tang and Si-Wu-Tang formulae. Sci. Rep. 6:34328. doi: 10.1038/srep34328

Tal, J., Piao, B., Samberg, L., Lazarov, N., and Sharf, M. (1985). Ultrasonographic and clinical correlates of metronidroz versus sequential clomiphene citrate: metronidroz therapy for induction of ovulation. Fertil. Steril. 44, 342–349. doi: 10.1016/S0015-0282(16)48858-8

Tseng, S. H., Lee, H. H., Chen, L. G., Wu, C. H., and Wang, C. C. (2006). Effects of three purgative decoctions on inflammatory mediators. J. Ethnopharmacol. 105, 118–124. doi: 10.1016/j.jep.2005.10.003

Wang, J., Qian, X., Gao, Q., Lv, C., Xu, J., Jin, H., et al. (2018). Quercetin increases the antioxidant capacity of the ovary in menopausal rats and in ovarian granulosa cell culture in vitro. J. Ovarian Res. 11, 51. doi: 10.1186/s13048-018-0421-0

Zhao, Z., Egashira, Y., and Sanada, H. (2003). Ferulic acid sugar esters are recovered in rat plasma and urine mainly as the sulfoglucuronide of ferulic acid. J. Nutr. 133, 1355–1361.

Zhao, Z., Egashira, Y., and Sanada, H. (2004). Ferulic acid is quickly absorbed from rat stomach as the free form and then conjugated mainly in liver. J. Nutr. 134, 3083–3088.

Zduńska, K., Dana, A., Kolodziejczak, A., and Rotsztejn, H. (2018). Antioxidant properties of ferulic acid and its possible application. Skin Pharmacol. Physiol. 31, 332–336. doi: 10.1159/000491755

Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2020 Lee, Kapelenier, Tsai, Lee, Xu and Wang. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.