Effects of Ethanol Extract from Sunkist \textit{(Citrus sinensis L. Osbeck)} Peels on Rat's Estrus Cycle

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Abstract

Kuhnle et al. has reported that Sunkist peels contain 36 µg/100 g wet weight of phytoesterogen. This study was designed to explore Sunkist peels' potential to recover the infertility due to ovulation failure. This study was performed in June 2020 at the Pharmacology Laboratory, Universitas Prima Indonesia, on 25 rats. These rats were divided into five groups: Control, Standard, and Sunkist Peels Extract-I, II, and III that received 1 ml of 0.5\% Sodium Carboxymethyl cellulose suspension, ethinyl estradiol 50 g/kgBB, 300 mg/kgBB, 450 mg/kgBB, and 600 mg/kg BW of ethanol extract from Sunkist peel, respectively. Every 8–12 hours, the rat's vagina were swabbed and smeared into an object glass to be stained by 10\% Giemsa and viewed at 40x and 100x magnifications. The leucocytes, especially neutrophils, epithelial cells, cornified and epithelial cells as well as the density of cells in vaginal swab were evaluated to determine the estrus cycle of the rats. The length of estrus phases was expressed as median (IQR) and analyzed using the Kruskall-Wallis test. The result of this study showed that the standard and highest dosage extract showed a significant extending of estrus and metestrus phases compared to the control group and significantly shortened diestrus phases. Hence, the highest dosage of Sunkist extract can prevent infertility by increasing the period of estrus cycle (ovulation) at the same level as the standard group.

Keywords: Estrus, ethinyl estradiol, Sunkist peels, vaginal swab

Pengaruh Ekstrak Etanol Kulit \textit{Sunkist} terhadap Siklus Estrus Tikus

Abstrak

Kuhnle et al. telah melaporkan bahwa kulit \textit{Sunkist} mengandung fitoesterogen sebesar 36 µg/100g berat basah. Penelitian ini dirancang untuk mengeksporasi potensi kulit \textit{Sunkist} untuk memulihkan infertilitas akibat kegagalan ovulasi. Penelitian ini dilakukan pada Juni 2020 di Laboratorium Farmakologi, Universitas Prima Indonesia dan menggunakan 25 ekor tikus yang dibagi atas lima kelompok, yaitu: kontrol, standar, ekstra kulit \textit{sunkist}-I, II, dan III yang menerima 1 mL suspensi 0,5\% \textit{Sodium Carboxymethyl cellulose}, etinilestradiol 50 µg/kgBB, 300 mg/kgBB, 450 mg/kgBB, dan 600 mg/kgBB ekstrak etanol kulit \textit{sunkist}. Setiap 8–12 jam, vagina tikus diusap dan dioleskan ke kaca objek. Kemudian, usapan vagina diwarnai dengan Giemsa 10\% dan dilihat pada pembesaran 40x dan 100x. Leukosit terutama neutrofil, sel epitel, sel epitel terkornifikasi, dan densitas sel pada usap vagina dievaluasi untuk mengetahui siklus estrus tikus. Panjang fase estrus dinyatakan sebagai median (IQR) dan dianalisis dengan \textit{kruskall-wallis}. Hasil penelitian menunjukkan bahwa ekstrak dosis standar dan dosis tertinggi menunjukkan perpanjangan fase estrus dan metestrus yang signifikan dibanding dengan kelompok kontrol dan memendekkan fase diestrus secara signifikan. Oleh karena itu, ekstrak \textit{sunkist} dosis tertinggi dapat mencegah kemandulan dengan meningkatkan periode siklus estrus (ovulasi) sebaik kelompok standar.

Kata kunci: Etil estradiol, estrus, kulit \textit{sunkist}, usapan vagina

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**Introduction**

During puberty, primary oocytes in primary follicles in the ovary undergo meiosis to form polar bodies and an ovum that will be released for conception during ovulation. Typically, the ovarium contains 300,000 reserve follicles. However, only around 400 follicles become mature and can be ovulated, while the remaining 99.7% will degrade.\(^1\) Ovulation is affected by various hormones, such as FSH (Follicle-Stimulating Hormone), LH (Luteinizing Hormone), estrogen, and progesterone. Before ovulation, estrogen takes a dominant role, while progesterone is dominant after ovulation. Normally, the estrogen level before menopause or premenopause is 30–400 pg/mL and is reduced at postmenopausal age to 0–30 pg/mL.\(^1,2\)

Infertility is a typical case found globally. Some reports show that the prevalence of infertility ranged from 9–18% among the general population. On the other hand, Novakovic et al. reported that more than 7 million individuals have conceived by Assisted Reproductive Technologies (ART) from 1978 until 2019.\(^3,4\)

Around 15% (48.5 million cases) of couples worldwide have infertility problems, where 17% of them come from industrial countries. The causes of infertility are, among others, ovulation failure, uterine tube damage, endometriosis, and a lower number in spermatozoa.\(^5,6\) In Indonesia, the Central Bureau for Statistics, Statistics Indonesia, reported that there are around 39.8 million women in childbearing age with 10–15% of them are infertile.\(^7\)

Various factors, both from the male and female sides, can cause infertility; however, more than 70% of problems are seen in females. Infertility in females can be caused by ovarium, uterine tube, uterine, hypothalamic, hypophysis, or systemic diseases. Around 27% of the cases are due to ovulation failure, both congenital and due to imbalanced hormones.\(^6,8,9\)

Indonesia is rich in various herbs, including some herbs that may contain phytoestrogens. Phytoestrogen is a natural compound with a similar structure to 17-β-Estradiol (E2) and a common sexual hormone in females. The phytoestrogens are commonly found as isoflavone, prenylflavonoid, coumestans, and lignans.\(^9\) One of the natural products containing these compounds is Sunkist (Citrus sinensis L. Osbeck) peels. The ethanol extract of Sunkist contains various phytochemicals, including a flavonoid which derivatives may act as the phytoestrogen. Due to its similar structure to the 17-β-Estradiol (E2), phytoestrogen affects the ovulation cycle. Therefore, the high intake of phytoestrogen may make the hypophysis increases the secretion of LH and FSH, which subsequently induces ovulation. Since one of the infertility causes is ovulation failure, phytoestrogens from the Sunkist peel may modify the ovulation cycle and prevent ovulation failure. Thus, this study was designed to explore the potential of Sunkist peels to prevent infertility caused by ovulation failure.\(^10,11\)

**Methods**

This was an experimental study performed in June 2020 at the Pharmacology Laboratory of the Faculty of Medicine, Universitas Prima Indonesia. This study was approved by the Health Research Ethics Committee of Universitas Prima Indonesia under the registration no. 012/KEPK/UNPRI/V/2020.

The materials used in this study were 96% ethanol, Sunkist peels, sodium carboxymethylcellulose (Na-CMC), Ethinyl Estradiol (Lynoral\(^\text{®}\)), aqua dest, rat pellets, chloroform, Giemsa staining (Catalogue Number: 109204), and alcohol. Female Wistar aged 3–6 months and weighed 200–220 grams were used as the animal models. The sample size was estimated by the Federer formulation, with a requirement that n should be higher than 4.75. Therefore, this study was performed on 25 Wistar rats, which were divided into five groups: control, standard, Sunkist Peels Extract-I, II, and III. All rats were acclimatized for two weeks before receiving the intervention.

The Sunkists used in this study were obtained from one of the traditional markets in Medan City and were identified at the Medanese Herbarium at The Faculty of Math and Science of University of North Sumatra. Five hundred grams of Sunkist peel was dried by aeration for seven days and then pulverized into the simplicia powder form.

The simplicia powder was extracted using the maceration method. Firstly, the powder was macerated by 70% ethanol as a solvent in a ratio of 1:10 for three days. The mixture was stirred regularly every day. After three days, the mixture was filtered by filter paper and the residue was then re-macerated with 600 mL of 70% ethanol for three days. The filtered re-macerated and macerated powder was evaporated by rotary evaporator at a temperature of 40°C and then concentrated using an oven at 40°C until it became thick extract.\(^12,13\)
The ethanol extract of Sunkist peels was then suspended into the sodium carboxymethylcellulose to form an oral suspension. The oral suspension used 0.5% Sodium Carboxymethyl cellulose as a vechiculum, which was made by mixing 0.5 grams Sodium Carboxymethyl cellulose powder into 30 mL hot aquadest in a mortar for 15 minutes until it formed a clear phase. It was then grounded to homogeny and dissolved in a volumetric flask by the remaining aqua dest. This suspension was also used by the vechiculum for extract and standard drugs.12

The ethinyl estradiol and Sunkist peels extract were suspended into the vehiculum in the next step, with 50 μg of ethinyl estradiol was suspended into the 5 mL vehiculum to form the ethinyl estradiol suspension. Meanwhile, the 300 mg, 450 mg, and 600 mg of Sunkist peels extract were suspended into 5 mL of vehiculum to form an oral suspension of Sunkist orange peel extract at a dose of 300 mg/kgBW, 450 mg/kgBW, and 600 mg/kgBW, respectively.14 These oral suspensions were given to each rat group.

Every 8–12 hours, the rats’ vaginals were swabbed by rotating a cotton swab that had been soaked into normal saline to the vaginal wall 360°. These vaginal swabs were smeared on a glass slide, and the slide was dried and fixed with methanol for 5 minutes, and stained with 10% Giemsa for 30 minutes. The preparations were washed carefully with running water and then dried. The slides were viewed by a microscope at 100x and 400x magnification.15,16

The vaginal swabs were evaluated based on the presence of leucocytes, especially neutrophils, epithelial cells, cornified epithelial cells, and the density of cells. The estrus phase was classified as proestrus, estrus, metestrus, and diestrus, based on the components.15,16

The length of each phase of the estrus cycle was shown as median (range) and analyzed by Kruskal-Wallis and Mann-Whitney. On the other hand, the total length of estrus cycles was shown as mean±SD and analyzed by one-way ANOVA and Post-Hoc Test Games Howell.

### Results

Vaginal swabs were obtained from each rat after 14 days of treatment and the microscopic view of some vaginal swabs were depicted in Figure 1 whereas the length of each phase and the overall cycle were shown in Table 1.

Based on Table 1, the provision of the standard and highest dose of Sunkist extract led to a significant extended estrus and metestrus phases than the control group and significantly shortened diestrus phases. On the contrary, the other two lower dosage groups showed an extended estrus and metestrus phase, but they did not reach the standard group’s length of phase. They also showed a shortened diestrus, but they did not attain the standard group’s length of the diestrus phase. Furthermore, no significant difference was seen in the estrus cycle’s total length of each group when compared to the control group, except for the group that received the lower dose of Sunkist peel extract. As a comparison, the normal length of each estrus cycle was shown in Table 2.

### Table 1 Analysis of Length of Estrus Phases and Estrus Cycles in Treatment Group

| Groups                  | Cycle (Hours)* | Total Length of Cycle** |
|-------------------------|----------------|-------------------------|
|                         | Proestrus      | Estrus                  | Metestrus    | Diestrus    |                     |
| Control                 | 8.00 (1.00)    | 33.50 (12.00)           | 10.00 (1.00) | 48.00 (24.00)| 100.30±9.38         |
| Standard                | 7.00 (0.00)a   | 56.00 (12.00)a          | 12.00 (0.00)a| 16.00 (10.00)a| 90.60±5.23          |
| Sunkist Peels Extract-I (300 mg/kgBW) | 8.00 (1.00)b   | 30.00 (12.00)b          | 10.00 (1.00)b| 24.00 (24.00)b| 81.20±13.66          |
| Sunkist Peels Extract-II (450 mg/kgBW) | 8.00 (1.00)b   | 36.00 (32.00)b          | 12.00 (2.00)b| 24.00 (24.00)b| 88.90±18.00          |
| Sunkist Peels Extract-III (600 mg/kgBW) | 7.00 (0.00)    | 56.00 (12.00)a          | 12.00 (3.00)a| 16.00 (6.00)a | 91.70±5.33          |
| p-value                 | <0.05          | <0.05                   | <0.05        | <0.05       | 0.001               |

* p-value was obtained from Kruskall wallis and Mann-Whitney and data was expressed as median (Range); ** p-value was obtained from one way ANOVA and Post Hoc Test Games Howell and data was expressed as Mean±SD; “Significantly difference against control groups; "Significantly difference against standard group
Based on the values in Table 2, the proestrus length from all groups was shorter than the normal range and this pattern was similar for the diestrus length. The diestrus length was shorter than the normal range, except for the control group. On the contrary, the estrus and metestrus phases were shown to be longer than the normal range. The estrus phase from the rats was determined by evaluation of the microscopic view of the vaginal swab.

Figure 1 presented the serial of the estrus cycle in the animal model. Figure 1A, 1C, 1E, and 1G depict the microscopic view of the vaginal swab at 100x magnification. Meanwhile, the Figure viz. 1B, 1D, 1F, and 1H demonstrate the microscopic view of the vaginal swab at 400x magnification. The figure shows the serial estrus cycle that includes estrus, proestrus, metestrus, and diestrus. The determination of the estrus phase was based on the following criteria.

Based on Table 3, it was clear that the estrus phase from the vaginal swab in Figure 1 matched the criteria of each estrus cycle. The cells density was lower in the proestrus (Figure 1A and 1B) and diestrus (Figure 1G and 1H) phases. In contrast, the estrus (Figure 1C and 1D) and metestrus (Figure 1E and 1F) phases had a higher density of cells. Other than using the density, the estrus cycle can be determined by leucocyte count of the vaginal swab. The leucocyte count was higher in the metestrus and diestrus cycles, and lower in others. The leucocyte is marked as
the rounded cell with a size that is smaller than the epithelial cells.

**Discussion**

This study proved the potentials of the Sunkist peels to improve infertility by demonstrating not only a longer estrus phase but also by a shorter total length of the estrus cycle. The estrus cycle of the rats is considered as an analogy to the human menstrual cycle.

The rat group that received the highest dosage of Sunkist peels extract had a prolonged estrus phase compared to the standard group. The prolonged phase will increase the opportunity for conception for the rats. Moreover, the rats that received the highest dose of the extract also showed a shorter total length of the cycle, preventing the delay of the next ovulation process. Rat as rodent animal has an estrous cycle similar to the human menstrual cycle (ovarian and uterine cycles). The estrus phases include proestrus, estrus, metestrus, and diestrus that occur around 4 to 5 days. The rats’ first estrous cycle begins after the pulsatile release of Luteinizing hormone (LH) after the fourth postnatal week (about 30 days). Afterwards, the rats are in puberty and, after hitting the first puberty, the rats are within anestrus phases for 8 to 9 days before entering their first proestrus, estrus, metestrus, and diestrus.\(^1\)

The rat estrous cycle resembles the human menstrual cycle. The proestrus phase of the rat corresponds to the human follicular stage. This phase is affected by the increasing estradiol and a little surge of prolactin that leads to the increase of LH and Follicle Stimulating Hormone (FSH). When the FSH reaches the peak concentration, it will cause a rapid decline of the estradiol level. Subsequently, the ovulation occurs based on the rat’s estrus phase. Consequently, the progesterone hormone becomes dominant, leading to the metestrus and diestrus phase. These cycles are reassembled in early and late human secretory stages, respectively.\(^1\)

This study show that the estrus phase in the estrous cycle of the group receiving the highest dose of Sunkist extract is longer. The lengthening of the estrus cycle in the highest extract dose group is due to the presence of phytoestrogen in the Sunkist peel extract. Phytoestrogen has a similar structure to endogen estradiol and binds with the alpha and beta receptors of estrogen. This complex will influence the gene expression of the cells. Kuhnle et al. reported that orange peels contain 36 µg/100g wet weight of phytoestrogen based on the LC-MS and subC-Labelled standard.\(^1\)

Also, Shalaby et al. reported that the presence of flavonones, flavones, and flavonols in *Citrus sinensis* and *Citrus aurantifolia* leaf or peel leads to a powerful estrogen-like activities. When the phytoestrogen affects the ovary, it will affect the oocyte maturation and interfere with the estrus cycle. The extension becomes important because it can reduce the cumulative number of cycles and become very important in the fertility cycle.\(^1,19,20\)

Overall, it can be concluded that the highest dose of Sunkist extract of 600 mg/kgBW could prevent infertility by increasing the period of the estrus cycle (ovulation).

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