Research Article

The Effect of $17\beta$-Estradiol Administration on Cutaneous Wound Healing in 24-Week Ovariectomized Female Mice

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Estrogen replacement promotes cutaneous wound healing in 8–10-week young ovariectomized female mice. However, research using aged ovariectomized female mice has not been reported, to the best of our knowledge. Therefore, we investigated the effect of $17\beta$-estradiol on cutaneous wound healing using 24-week middle-aged ovariectomized female mice. Twenty-week-old female mice were divided into three groups: medication with $17\beta$-estradiol after ovariectomy (OVX + $17\beta$-estradiol), ovariectomy (OVX), and sham (SHAM). After 4 weeks, the mice received two full-thickness wounds. Then, the OVX + $17\beta$-estradiol group was administered $17\beta$-estradiol at 0.01 g/day until healing. The ratio of wound area in the OVX + $17\beta$-estradiol group was significantly decreased compared with that in the OVX group. The numbers of neutrophils and macrophages in the OVX + $17\beta$-estradiol group were significantly smaller than those in the OVX group. In addition, the ratio of myofibroblasts in the OVX + $17\beta$-estradiol group was significantly higher than that in the OVX group. These data suggested that exogenous continuous $17\beta$-estradiol administration promotes cutaneous wound healing in 24-week OVX female mice by reducing wound area, shortening inflammatory response, and promoting wound contraction. However, it is unclear whether the effect of exogenous estrogen on wound healing outweighs the delay of wound healing due to advanced age.

1. Introduction

Cutaneous wound healing is a complex, tightly orchestrated response to injury, carefully regulated at temporal and spatial levels [1]. Generally, there are various overlapping phases of the repair process (the immediate response; the inflammatory response; the proliferation, migration, and contraction phase; and the remodeling phase) [1].

With advanced age, this series of events becomes disrupted and healing is delayed. In elderly humans, delayed healing of acute wounds is characterized by the extension of inflammation [2], increased protease activity [3, 4], and delayed re-epithelialization [5]. However, postmenopausal women with systemically reduced estrogen heal poorly [5], whereas exogenous estrogen treatment reverses this delayed cutaneous wound healing by dampening excessive neutrophil recruitment [6], promoting re-epithelialization [5] and increasing collagen deposition [5–7].

Rodent models subjected to ovariectomy (OVX) constitute a cornerstone in elucidation of the effects and detailed biological mechanisms of estrogen. OVX exhibited delayed cutaneous wound healing compared with SHAM [5, 6, 8–14], whereas exogenous estrogen treatment reversed this delay by decreasing wound area [5, 8, 12–14], local numbers of neutrophils and macrophages [8–12, 14], and wound levels of TNFα [8–12], and by promoting re-epithelialization [5, 9, 11–14], and collagen deposition [5]. However, these studies evaluating the effects of estrogen used 8–10-week young female mice, and research using aged female mice has not been reported, to the best of our knowledge.

Since the relationship between advanced age and exogenous estrogen treatment in relation to cutaneous wound
healing remains unknown, we here attempt to verify the effect of estrogen on cutaneous wound healing using an older OVX female mouse model. We investigate the effect of 17β-estradiol on cutaneous wound healing using 24-week middle-aged OVX female mice. The aims of the present study are to evaluate how 17β-estradiol administration promotes cutaneous wound healing in 24-week middle-aged OVX female mice, and which factor has a stronger influence on cutaneous wound healing: estrogen or age.

2. Materials and Methods

2.1. Animals. Sixty-three C57BL/6 female mice aged 8 weeks (Sankyo Lab Service Co., Tokyo, Japan) were used in the experiments. They were caged individually in an air-conditioned room at 25.0 ± 2.0°C with light from 08:45 to 20:45 hours, and water and chow were given freely. The experimental protocol and animal care were in accordance with the Guidelines for the Care and Use of Laboratory Animals of Kanazawa University, Japan (AP-122316).

2.2. Wounding. The mice were bred until 20 weeks. At this time, they were anesthetized by intraperitoneal (IP) injection of pentobarbital sodium (0.05 mg/g weight), and the dorsum was shaved. Then, they were subjected to sham surgery (SHAM) or ovariectomy (OVX) according to OECD guidelines [15]. Four weeks later, they were divided into three groups (21 mice/group): SHAM group, OVX group, and OVX + 17β-estradiol group. Then, under anesthesia with shaving, two circular full-thickness skin wounds (4 mm in diameter) including the panniculus carnosus muscle on both sides of the dorsum of the mouse were made with a Kai sterile disposable biopsy punch (Kai Industries Co. Ltd., Gifu, Japan). In SHAM and OVX groups, the wounds were covered with hydrocolloid dressing (Tegaderm; 3M Health Care, Tokyo, Japan) to maintain a moist environment, and then the mouse was wrapped with sticky bandages (Meshpore Tape; Nichiban, Tokyo, Japan). They were changed every day. In the OVX + 17β-estradiol group, wounds received the same treatment. However, after wound treatment, they were also treated with 0.01 g of 17β-estradiol gel (Lestrogel 0.06%; Bayer Yakuhin, Osaka, Japan). It was placed on clean gauze using a 1-mL syringe and applied to the skin on the back avoiding the wounds every day.

2.3. Macroscopic Observation. The day when wounds were made was designated as day 0, and the process of wound healing was observed from days 0 to 14 after wounding. Wound edges were traced on polypropylene sheets and photographs were taken every day. The traces on the sheets were captured with a scanner onto a personal computer using Adobe Photoshop Elements 7.0 (Adobe System Inc., Tokyo, Japan), and the areas of wounds were calculated using image analysis software Scion Image Beta 4.02 (Scion Corporation, Frederick, Maryland, USA). Wound area is shown as the ratio of wound area every day to the initial wound area on day 0 when the wound was made. Our previous studies reported [16–18] that a full-thickness cutaneous wound on the back of a mouse without any management, such as the application of honey, daidzein, or estrogen, healed with scar formation on about day 14 after wounding and the ratio of wound area reached about 0.1. Therefore, we calculated the number of wounds reaching the 0.1 ratio of wound area per group.

2.4. Plasma 17β-Estradiol and Uterus Assay. The mice were euthanized by a massive pentobarbital sodium IP injection on day 14. Plasma was prepared from each mouse’s blood isolated through cardiac puncture and frozen until the time of assay. Plasma 17β-estradiol levels were determined by radioimmunoassay (RIA), and since levels less than 10 pg/mL could not be detected, such levels were recorded as 10 pg/mL. This was outsourced to the manufacturer of this assay (Mitsubishi Chemical Medience Corporation, Tokyo, Japan). The uterus was harvested according to OECD guidelines [15] after blood isolation, its wet weight was measured, and it was fixed in 4% paraformaldehyde and embedded in paraffin.

2.5. Plasma TNF-α. The mice were euthanized by a massive pentobarbital sodium IP injection on days 3 and 7. Plasma was prepared from each mouse’s blood isolated through cardiac puncture and frozen until the time of assay. Plasma TNF-α levels were determined by ELISA (R&D Systems, Tokyo, Japan) according to the manufacturer’s guidelines.

2.6. Histological Procedure and Immunohistological Staining. The mice were euthanized by a massive pentobarbital sodium IP injection on days 3, 7, 11, and 14 after wounding. The wound and the surrounding intact skin were harvested and each sample of wound and surrounding intact skin was bisected at the wound center. One-half of each wound was stapled onto polypropylene sheets to prevent overcontraction of the sample and fixed in 4% paraformaldehyde for 12 hours. These samples were dehydrated in an alcohol series, cleaned in xylene, and embedded in paraffin to prepare 5-μm serial paraffin sections. The remainder of each wound was embedded in tissue-Tek OCT (Sakura Finetek, Japan) before fixing to prepare 5-μm serial ice sections. At least 6 serial sections near the center of the wound were obtained from one wound and stained according to the following methods. Paraffin sections of 5-μm thickness were stained with hematoxylin and eosin (H&E) or subjected to Azan staining and immunohistologically stained with anti-neutrophil antibody at a concentration of 1:100 (Abcam Japan, Tokyo, Japan) for detecting neutrophils, anti-Mac-3 antibody at a concentration of 1:100 (BD Pharamingen, Tokyo, Japan) for detecting macrophages, and cryosections of 5-μm thickness were immunohistologically stained with anti-α-smooth muscle actin (α-SMA) antibody at a concentration of 1:500 (Abcam Japan, Tokyo, Japan) for detecting myofibroblasts. Negative control slides were obtained by omitting each primary antibody.

2.7. Microscopic Observations. Images were imported onto a computer using a digital microscopic camera (DP2-BSW Olympus, Japan). Measurements for the proportions lacking...
re-epithelialization were performed using DP2-BSW Olympus software: the distance between both wound edges and the distance between the tips of elongated new epithelium were measured, and then the latter was divided by the former (no re-epithelialization length/wound length). Measurements for collagen deposition colored blue (collagen pixels/total wound pixels) and for myofibroblasts colored brown (myofibroblast pixels/total wound pixels) were performed using Adobe Photoshop Elements 7.0 as follows: the wound area was first selected; one wound edge, the surface of the wound, the other wound edge, and the bottom of the wound, which is the position of the panniculus carnosus, were surrounded, and then the number of pixels in the surrounded area (=wound area) was calculated. Next, the collagen deposition colored blue or myofibroblasts colored brown were selected and the number of pixels of the blue or brown area (the area of collagen deposition or the area of myofibroblasts) was calculated; finally, the number of pixels of the area of collagen deposition or the number of pixels of the area of myofibroblasts was divided by the number of pixels of the wound area. To analyze the numbers of neutrophils and macrophages in the granulation tissue, each positive cell was counted by observation through a light microscope using a ×40 objective at three sites of granulation tissue: two sites near the two wound edges and the center of the granulation tissue. Areas of these three sites were calculated on the monitor of the DP2-BSW and the total number of neutrophils or macrophages at the three sites was divided by the whole area of these three sites.

2.8. Statistical Analysis. Data are expressed as mean ± SD and analyzed using JMP 8.0.1 (SAS, USA). Fisher’s exact probability test, ANOVA, and Student’s t-test or Tukey-Kramer multiple comparison test was performed. Differences were considered significant at $P < 0.05$.

3. Results

3.1. Uterine Weight and Plasma 17β-Estradiol Levels. We confirmed that the ovaries had been removed successfully in the OVX and OVX + 17β-estradiol groups. The body weight on day 14 after wounding was 25.08 ± 0.90 g in the SHAM group, 25.45 ± 0.38 g in the OVX group, and 25.33 ± 2.67 g in the OVX + 17β-estradiol group. The macroscopic and histological features of the uterus are shown in Figures 1(a) and 1(b). The uterine weight in the OVX + 17β-estradiol group was significantly greater than the uterine weights of SHAM and OVX groups on day 14 ($P < 0.01$). There were no significant differences between SHAM and OVX groups on day 14 (Table 1). The plasma 17β-estradiol level in the OVX + 17β-estradiol group was significantly greater than those levels of SHAM and OVX groups on day 14 ($P < 0.01$). There were no significant differences between SHAM and OVX groups on day 14 (Table 1), although in the OVX group, 4 out of 6 mice had plasma 17β-estradiol levels under 10 pg/mL and, in the SHAM group, 1 out of 6 mice did. Estrus and proestrus were not observed in the SHAM group when we observed smears of the SHAM mice before euthanizing them.
Table 1: Plasma 17β-estradiol levels and uterine weight.

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<th>SHAM</th>
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<th>OVX + 17β-estradiol</th>
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<td>17β-estradiol (pg/mL)</td>
<td>11.3 ± 1.0**</td>
<td>10.8 ± 1.8**</td>
<td>27.7 ± 7.7</td>
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<tr>
<td>Uterine weight (mg)</td>
<td>35.0 ± 9.0**</td>
<td>34.0 ± 16.0**</td>
<td>122.0 ± 21.0</td>
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Values are expressed as mean ± SD, n = 6 for each group, ANOVA, Tukey-Kramer. **P < 0.01 with respect to the OVX + 17β-estradiol group.

3.2. Wound Area. In the SHAM group, wound areas increased for 2 days and then decreased rapidly until day 8, after which they decreased slowly until day 14 (0.15 ± 0.08, ratio of wound area to initial wound area on day 14). The ratio of wound area to initial wound area of 5 out of 12 wounds reached a ratio of 0.1 to the initial wound area on day 14. In the OVX group, wound areas increased for 2 days and then decreased rapidly until day 9, after which they decreased slowly until day 14 (0.24 ± 0.15). The ratio of wound area to initial wound area of none out of 12 wounds reached a ratio of 0.1 on day 14. On the other hand, in the OVX + 17β-estradiol group, wound area increased only for 1 day and then decreased rapidly until day 11, after which it decreased slowly until day 14 (0.08 ± 0.03). The ratio of wound area to initial wound area of 9 out of 12 wounds reached a ratio of 0.1 on day 14. The ratio of wound area in the OVX + 17β-estradiol group was significantly smaller than that of the OVX group on days 3 and 7, with a significant decrease from days 3 to 7 in the OVX + 17β-estradiol group (P = 0.0137 and 0.0145, resp.). There were no significant differences of the ratio of wound area between OVX + 17β-estradiol and SHAM groups, or between SHAM and OVX groups on days 0–14. However, the mean ratio of wound area in the OVX group was larger throughout the whole period, except for the first day (Figure 2(b)). The number of wounds reaching a ratio of wound area of 0.1 on day 14 after wounding was significantly different between SHAM and OVX groups and between OVX and OVX + 17β-estradiol groups (P = 0.0373 and 0.0003, resp.), but there were no significant differences between SHAM and OVX + 17β-estradiol groups.

3.3. Neutrophils, Macrophages and Plasma TNF-α Levels. The number of neutrophils in the OVX + 17β-estradiol group was significantly smaller than that in the OVX group on day 3 (P = 0.0326). There were no significant differences between OVX + 17β-estradiol and SHAM groups or between SHAM and OVX groups on day 3, and there were no significant differences among the three groups on day 7 (Figure 3(a)). The numbers of neutrophils significantly decreased from days 3 to 7 in the OVX and OVX + 17β-estradiol groups (P = 0.0137 and 0.0145, resp.).

The number of macrophages in the OVX + 17β-estradiol group was significantly smaller than that in the OVX group on day 3 (P = 0.0012). There were no significant differences between OVX + 17β-estradiol and SHAM groups, or between SHAM and OVX groups on day 3, and there were no significant differences among the three groups on day 7 (Figure 3(c)). The number of macrophages decreased in the OVX group from days 3 to 7 but remained unchanged in the SHAM and OVX + 17β-estradiol groups.

There were no significant differences of plasma TNF-α levels among the three groups on days 3 and 7 (Figure 3(e)).

Figure 2: Macroscopic wound healing. (a) Four mm diameter wounds were produced and healing was recorded by photography. Bar 5 mm. (b) The ratios of wound areas to initial area on day 0 are shown on line graphs for each day. There were significant differences between the OVX + 17β-estradiol and OVX groups on days 4 and 11–14. However, there were no significant differences between the SHAM and OVX groups or between the SHAM and OVX + 17β-estradiol groups. Values are expressed as mean ± SD, n = 12, for each group, ANOVA, Tukey-Kramer. *P < 0.05; OVX versus OVX + 17β-estradiol.

Plasma TNF-α levels in the SHAM and OVX + 17β-estradiol groups decreased from days 3 to 7, with a significant decrease in the OVX + 17β-estradiol group (P = 0.0233), but it remained unchanged in the OVX group.

3.4. Re-Epithelialization, Collagen Deposition, and Wound Contraction. On day 3, new epithelium appeared from the
Figure 3: Inflammatory cells and plasma TNF-α levels. (a) The number of neutrophils per mm². The number of neutrophils in the OVX + 17β-estradiol group was significantly smaller than that in the OVX group on day 3. (b) Neutrophils (arrows) stained with antineutrophil antibody were observed in wound tissue on days 3 and 7. Bar 20 μm. (c) The number of macrophages per mm². The number of macrophages in the OVX + 17β-estradiol group was significantly smaller than that in the OVX group on day 3. (d) Macrophages (arrows) stained with anti-Mac-3 antibody were observed in wound tissue on days 3 and 7. Bar 20 μm. (e) Systemic TNF-α levels, determined by ELISA on serum samples. There were no significant differences between the three groups on days 3 and 7. Values are expressed as mean ± SD, n = 4-5 for each group, ANOVA, t-test, or Tukey-Kramer; *P < 0.05, **P < 0.01: OVX versus OVX + 17β-estradiol.

wound edge. It gradually covered the wound surface along with wound healing. By day 11, new epithelium completely covered the wound surface in the OVX + 17β-estradiol group, but 1 out of 5 wounds was not completely covered in the SHAM and OVX groups. By day 14, the surfaces of all wounds in the three groups were covered with new epithelium. However, there were no significant differences among the three groups on days 3–14 (Figure 4(a)).

Collagen deposition became thick and had a high density with the progression of wound healing in the three groups. There were no significant differences among the three groups on days 7–14, but that in the OVX + 17β-estradiol group tended to be larger than that in the OVX group on days 7 and 11 (P = 0.0648 and 0.0799, resp.) (Figure 4(c)).

A few myofibroblasts were observed at the wound site in the three groups on day 7. By day 11, many myofibroblasts
Figure 4: Re-epithelialization, collagen deposition, and wound contraction. (a) Proportion of no re-epithelialization. There were no significant differences between the three groups on days 3–14. (b) Hematoxylin and eosin sections on day 7. Bar 500 μm. (c) The ratio of collagen fibers. There were no significant differences between the three groups, although that in the OVX + 17β-estradiol group tended to be increased compared with that in the OVX group on days 7 and 11. (d) Azan-stained sections on day 7. Bar 500 μm. (e) The ratio of myofibroblasts. The ratio of myofibroblasts in the OVX + 17β-estradiol group increased significantly compared with that in the OVX group on day 11. However, there were no significant differences between OVX + 17β-estradiol and SHAM groups or SHAM and OVX groups. (f) Myofibroblasts stained with anti-α-SMA antibody were observed in granulation tissue on day 11. Bars 500 μm (SHAM), 1000 μm (OVX), and 200 μm (OVX + 17β-estradiol). Note a bridge-like structure (arrows) formed by the myofibroblasts. Values are expressed as mean ± SD, n = 4–5 for each group, ANOVA, Tukey-Kramer; *P < 0.05: OVX versus OVX + 17β-estradiol.
were observed in the granulation tissue in the three groups, building bridge-like structures. They had almost completely disappeared by day 14, with no detection in the three groups. The ratio of myofibroblasts in the OVX + 17β-estradiol group was significantly larger than that in the OVX group on day 11 ($P = 0.0426$). However, there were no significant differences between OVX + 17β-estradiol and SHAM groups or between SHAM and OVX groups on days 7–14 (Figure 4(e)).

4. Discussion

In the present study, the ratio of wound area in the OVX + 17β-estradiol group was significantly smaller than that in the OVX group on days 4 and 11–14 after wounding. This shows that exogenous, continuous 17β-estradiol administration is effective for decreasing wound area in 20-week OVX female mice. This agrees with previous research that cutaneous wound healing in OVX mice is promoted by exogenous estrogen treatment via a decrease of wound area [5, 8, 12–14]. On the other hand, in the present study, the ratio of wound area was not significantly different between SHAM and OVX + 17β-estradiol groups. The same result can be assumed from graphs in previous research papers, although they do not clearly show that cutaneous wound healing in SHAM mice was not delayed compared with that in OVX mice administered 17β-estradiol at the age of 8–10 weeks [9–11]. Moreover, in the present study, the ratio of wound area was also not significantly different between SHAM and OVX groups. This contradicts previous research that cutaneous wound healing in OVX mice was delayed compared with that in SHAM mice [5, 6, 8–14]. However, the number of wounds reaching a ratio of wound area of 0.1 on day 14 after wounding in the present study in the SHAM group was significantly larger than that in the OVX group. This shows that cutaneous wound healing in SHAM mice is promoted with that in OVX mice by physiological estrogen. From these findings, it is assumed that cutaneous wound healing in OVX mice is the most delayed, and cutaneous wound healing in OVX mice with exogenous, continuous 17β-estradiol administration is the most promoted, while cutaneous wound healing in SHAM mice is intermediate between these two groups. These findings indicate that exogenous, continuous 17β-estradiol administration is more effective for cutaneous wound healing than the natural physiological 17β-estradiol that fluctuates with estrus [19].

These differences in cutaneous wound healing among OVX, SHAM, and OVX + 17β-estradiol groups in the present study cannot necessarily be interpreted by the present results: the ratio of re-epithelialization, the numbers of neutrophils and macrophages, the level of TNF-α, and the ratios of myofibroblasts and collagen deposition. Previous researchers reported that the physiological level of 17β-estradiol and exogenous, continuous 17β-estradiol administration after OVX promoted cutaneous wound healing by promoting re-epithelialization [5, 9, 11–14], decreasing local numbers of neutrophils and macrophages [8–12, 14] as well as wound level of TNFα [8–12], and promoting collagen deposition [5]. In addition, Emmerson et al. reported that smooth muscle action (SMA) was reduced in day 7 wound tissue from OVX mice compared with that of wild-type mice, which have the physiological level of 17β-estradiol [13]. On the other hand, in the present study, although there were significant differences only between OVX and OVX + 17β-estradiol groups—the numbers of neutrophils and macrophages in the OVX + 17β-estradiol group were significantly smaller than those in the OVX group on day 3, and the ratio of myofibroblasts in the OVX + 17β-estradiol group increased significantly more than that of the OVX group on day 11—there were no significant differences in the ratios of collagen deposition and re-epithelialization or the level of TNF-α between OVX and OVX + 17β-estradiol groups. However, there were no significant differences of these factors between OVX and SHAM groups. These findings may indicate that exogenous, continuous 17β-estradiol administration is more effective than the natural physiological estrogen that fluctuates over the course of an estrous cycle, as we have discussed for the wound area.

However, from these findings, the following question arises. Why were there no significant differences between OVX and SHAM mice? Previous research evaluated the effects of estrogen on cutaneous wound healing using 8–10-week young female mice in comparison among OVX, SHAM, and OVX with exogenous 17β-estradiol administration. In the present study, in contrast to previous research, we used aged 24-week mice. Mills et al. demonstrated that cutaneous wound healing in old male mice was delayed by extending the wound area, compared with that in young male mice [20]. Hardman and Ashcroft reported, from a microarray study, that 78% of differentially expressed genes were estrogen-regulated, while only 3% were age-associated [21]. The results in the present study suggest that, although exogenous, continuous 17β-estradiol administration influences cutaneous wound healing, the influence of age cannot be disregarded since the effect of the SHAM group with physiological estrogen on cutaneous wound healing seems to be intermediate between the effects of the OVX and OVX + 17β-estradiol groups. Therefore, we will conduct further research using an older mouse model and attempt to evaluate the influence of 17β-estradiol and advanced age on cutaneous wound healing in the near future.

5. Conclusions

We clarified that exogenous, continuous 17β-estradiol administration has beneficial effects on wound area, inflammatory cells, and myofibroblasts during cutaneous wound healing. Therefore, our research suggests that exogenous, continuous 17β-estradiol administration also promotes cutaneous wound healing in 24-week OVX female mice by reducing wound area, shortening inflammatory response, and promoting wound contraction. However, it is unclear whether the effect of exogenous, continuous 17β-estradiol administration on wound healing outweighs the delay of wound healing due to advanced age.
Conflict of Interests
The authors declare that there is no conflict of interests regarding the publication of this paper.

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