

Protective Effects of Micro-Injected Adipose-Derived Mesenchymal Stem Cells on Ovarian Ischemia/Reperfusion Injury in Rats

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Introduction: The authors investigated the immunohistochemical, histological, and biochemical effects of adipose-derived stem cells (ADSC) which possess antioxidant and anti-inflammatory properties in the ovarian ischemia/reperfusion rat model.

Materials and Methods: Rats were randomly allocated to sham, ADSC, torsion/detorsion, torsion/detorsion/ADSC groups. ADSCs were micro-injected to torsion/detorsion/ADSC during the detorsion procedure after a 6-h period of torsion. BrdU, caspase-3, TUNEL, eNOS, and iNOS staining were used. Advanced oxidation protein products (AOPPs), superoxide dismutase (SOD) activity, and levels of total sulfhydryl (TSH) were evaluated.

Results: Serum AOPP increased in the torsion/detorsion, tissue and serum AOPP levels in the torsion/detorsion/ADSC group were decreased. Tissue SOD activity in torsion/detorsion and torsion/detorsion/ADSC groups was significantly decreased compared to the sham group. The iNOS was stronger in the torsion/detorsion group following torsion and detorsion but decreased after ADSC microinjection in the torsion/detorsion/ADSC group. ADSC treatment after torsion-detorsion also reduced caspase-3 immunoreactivity and apoptotic cell number.

Conclusion: ADSC implementation after torsion/detorsion might protect the ovary from ischemia/reperfusion injury.

Keywords: Adipose-derived stem cells, Ischemia, Ovary, Rat, Torsion

Introduction

Torsion of the adnexa, the rotation of the fallopian tube and ovary, accounts for 2.7–3% of gynecological emergencies (1). Since most cases occur at reproductive ages, early diagnosis and effective treatment are vital for

maintaining the ovarian reserve and hormonal functions. Salpingo-oophorectomy was the previously preferred management option due to risks of thromboembolism and potential malignancy with a nonviable appearance of the

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ovary; but recent studies recommend ovarian detorsion for patients desiring to protect their fertility (2). In ovarian torsion, gangrene and necrosis with venous and lymphatic occlusion and subsequent arterial insufficiency may ultimately cause loss of ovarian reserve due to ischemia (3). After reperfusion of the ovary by detorsion, many reactive oxygen species (ROS), namely hydroxyl radicals, hydrogen peroxide, and superoxide radicals develop due to increased oxygen levels (4,5). The elevated ROS result in DNA damage, endothelial destruction, and apoptosis of granulosa cells (6,7). To improve the outcomes of ovarian detorsion operations, many promising anti-inflammatory and antioxidant agents have been investigated such as atorvastatin, *Galega officinalis*, *Olea europaea* and enoxaparin (8-11). Human adipose-derived stem cells (ADSCs) are derived from human adipose tissue as a kind of superior biomaterial that can be suitable for allotransplantation and regenerative medicine (12). ADSCs stably expand under basic culture conditions, exhibit a low risk of contamination, and they have low immune rejection following transplantation (13). ADSCs also possess antioxidant and anti-inflammatory properties and have recently been used in many rat I/R models in the liver and brain (12,14). Stem cells are also being investigated for obstetric and gynecological applications such as in Asherman syndrome, stress urinary incontinence, and premature ovarian failure (15-17). Furthermore, previous studies showed that cytokines secreted from ADSCs could resist the process of natural ovarian aging (18). Recently it has been demonstrated that human ADSC-derived exosomes recover the function of premature ovarian failure by increasing the proliferation rate and inhibiting the apoptosis rate of the human granulosa cells (19).

In this study, we aimed to investigate the effects of ADSCs in I/R injury caused by ovarian torsion/detorsion surgery.

Materials and Methods

Animals

A total of 25 adult, virgin, 10–12 weeks old female Wistar albino rats were allocated in this study. The animals (170–210 g) were housed under conditions of 20–24°C, 12-h of day/night cycles, and 50–60% humidity. The Dokuz Eylül University Ethics Committee for Animal Research approved the animal protocol for this study (Approval no. 28/2015). Animal research investigations conformed to the experimental guidelines in line with Dokuz Eylül University Committee for Human Care.

ADSC Isolation

ADSCs were isolated from five rats that were not allocated to experimental groups. Anesthesia was induced by intraperitoneal 10 mg/kg (Rompun, Bayer, Leverkusen, Germany) xylazine hydrochloride, and 50 mg/kg ketamine administration (Ketalar, Eczacıbaşı, Turkey). The antiseptic solution was used for surgical site preparation and the bilateral groin fat pads were removed. Adipose tissues were densely processed in sterile phosphate-buffered saline (PBS) with 5% penicillin-streptomycin.

Tissues were dissociated with scalpels and incubated for 30 min in α -MEM containing 0.1% collagenase Type 1 at 37°C in a 5% CO₂. Tissues were then transferred to media for culture (20% fetal bovine serum (FBS), α -MEM, 1% L-glutamine), and after being centrifuged twice, they were cultivated in humidified incubators in α -MEM/10% FBS, 50 μ g/ml streptomycin and 50 U/ml penicillin at 37°C and 5% CO₂ (20). The media were changed every two days during the 10-day cultivation period.

Characterization of ADSCs

The cells were fixed in 4% paraformaldehyde for 30 minutes in PBS and permeabilized for 15 min with 0.1 Triton X-100 (A4975, AppliChem, Darmstadt, Germany) after washing with PBS. Endogenous peroxidase activity of cells was put out by incubation with 3% hydrogen peroxide

(K31355100, Merck, Darmstadt, Germany) for 5 min at room temperature. Afterward, cells were washed with PBS. After washing, cells were incubated in blocking serum for 1h in compliance with the manufacturer's instructions (85-9043, Invitrogen, Camarillo, CA, USA). Anti-CD-34 (ab51037, Abcam) and anti-CD-44 (BA3382, BOSTER) primary antibodies were applied at a dilution of 1:25 and 4°C overnight (20). Primary antibodies were removed and cells were washed with PBS. Biotinylated goat IgG, anti-rabbit/mouse IgG secondary antibody, and peroxidase-conjugated streptavidin (D01-110, Life Science Division), were sequentially added, with each step incubated for 30 min.

Cells were washed three times with PBS between these incubation periods. Then, the samples were incubated with diaminobenzidine /hydrogen peroxide (00-2020, Invitrogen) for 5 min. Mayer's hematoxylin (HMM999, Scytek Laboratories, West Logan, Utah, USA) was used for counterstaining. The tissues were washed in distilled water, mounted with mounting medium (AML060, Scytek), and covered with coverslips for evaluation under the light microscope (Olympus BX40, Olympus Corp., Tokyo, Japan). The same procedures were applied for the control samples using the same type of IgGs instead of primary antibodies.

Treatment Groups

Animals were randomly distributed: group 1 (sham, saline n: 4), group 2 (ADSCs, n:7), group 3 (torsion + detorsion + saline, n:6), group 4 (torsion + detorsion + ADSCs, n:8).

Surgical Procedure

Rats underwent surgery in a supine position under general anesthesia as described in ADSC Isolation. After shaving and cleaning the surgical sites with antiseptic solution (Baticon, Drogsan, Turkey), 2.5-cm midline vertical incisions were made.

Group 1: Uterine corns and adnexa were visualized. Then, 20 µl saline was administered separately to the ovaries with a microinjector.

Group 2: After exposing the adnexa, 1×10^5 cells/ml ADSCs in 20 µl PBS (including 2000 cells of ADSCs) were separately administered to the ovaries with a microinjector (13,21).

Group 3: To create torsion, the adnexa with connecting vessels were rotated 720° clockwise, and rotated adnexa were affixed to the anterior wall by 5/0 silk sutures. The abdominal wall incision was closed. Following a 6-h period of torsion, a re-laparotomy was performed for detorsion (reperfusion) by rotating the ovaries 720° counterclockwise and both ovaries were exposed. Then, 20 µl saline was administered separately.

Group 4: 6-h torsion and detorsion procedures were performed as described above. After the ovaries were exposed in the detorsion procedure, 1×10^5 cells/ml ADSCs in 20 µl PBS (2000 cells of ADSCs) were administered separately with a microinjector (13,21).

Abdominal wall incisions were all closed with 3/0 silk sutures. After surgical recovery, animals were housed separately. Following 7 days of the recovery period, all animals underwent laparotomy, and the oophorectomies were performed. All procedures were performed by the same researchers. One of the paired ovaries was selected randomly and fixed in 10% buffered formalin solution for histological and immunohistochemical analysis, while the other one was packed in oiled foil paper for further biochemical analysis. Just before sacrifice of the rats via hypovolemic shock, 2 ml of blood was withdrawn from the inferior vena cava. After 15 min centrifugation of blood samples at 5000 rpm, the serum was collected in Eppendorf tubes. The Eppendorf tubes and packages of ovaries in oiled foil papers were transferred to dry ice and stored at -20°C until final biochemical analysis.

Histopathological Examination

Tissue samples were fixed for 48 hours in a 10% buffered formalin solution, dehydrated through alcohol, and inserted in paraffin. Paraffin blocks were sliced into sections with 5- μ m-thickness (Leica RM2125RTS, Leica Biosystems, Nussloch, Germany). Hematoxylin and eosin staining was used as a routine protocol. The sections were examined for the severity of ovarian damage, such as edema, inflammation, hemorrhage, and congestion. All sections were examined under light microscopy (Olympus BX40, Tokyo, Japan) using the image analysis method (Image-Pro Plus 5.1). Tissue sections were analyzed by the same experienced histologist.

Immunohistochemical Analysis

For immunohistochemical analysis, Terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL), Bromodeoxyuridine/5-Bromo-2'-deoxyuridine (BrdU), caspase-3, Endothelial nitric oxide synthase (eNOS), and inducible nitric oxide synthase (iNOS) stainings were used. The intensity of immunolabeling was graded as follows: weak (+); moderate (++); and strong (+++). This grading was performed by two observers who were blinded to the experimental conditions.

TUNEL Assay

The slides were mounted with Entellan® (Merck 107961.0500) and examined by two histologists independently. 100 TUNEL-negative or positive-cells were counted per case on randomly chosen fields. The TUNEL-positive cells' percentage was determined.

Biochemical Analysis

Ovarian tissues were homogenized in 20 mM phosphate buffer at pH 7.4. Homogenized samples were centrifuged at 14000 rpm at 4°C for 15 min. After centrifugation, supernatants were divided into aliquots for analysis of superoxide dismutase (SOD) activity, advanced oxidation protein products (AOPP), total sulfhydryl (TSH), and total protein levels, while

pellets were used for the myeloperoxidase (MPO) activity.

Measurement of SOD Activity

The SOD activity of ovarian tissues or serum was measured spectrophotometrically (22). SOD activity in ovarian and serum were given as units/mg protein and units/mL, respectively.

Assay of TSH

The TSH levels of ovarian tissue or serum were evaluated with a linear standard curve formed by using reduced glutathione (23). Protein levels in supernatants with sodium dodecyl sulfate were measured according to Lowry et al (24). TSH levels were expressed as μ mol/mg protein and μ mol/L for tissue or serum respectively.

Assay of AOPP Levels

AOPP levels in ovarian tissue and serum were measured according to the spectrophotometric method of Witko-Sarsat (25).

Assay of MPO Activity

Bradley's method was used for measuring MPO activity in the ovary (26).

Total Protein Assay

Total protein levels in ovarian tissue were measured according to the spectrophotometric method (27).

Statistical Analysis

The authors determined the sample size from the results obtained from the rats that they evaluated preliminarily. From the differences, a domineering two-tailed α value of 0.05 and a β value of 0.20 (study power: 80%). SPSS version 22.0 software (Chicago, IL, USA) was used. Distribution normality was assessed with the Shapiro-Wilk test. Results are given as mean $\pm$ standard error or median (min-maximum). Differences among experimental groups were determined by one-way analysis of variance (ANOVA) and Tukey's post hoc tests. Kruskal-Wallis was used for analyzing non-parametric data. $p < 0.05$ was considered significant.

Results

Results of ADSC Characterization

ADSCs were CD 34 (-) (negative) and CD 44 (+) (positive) (Fig.1). ADSCs were confirmed not to be hematopoietic stem cells as being CD34 (-) and confirmed to be adipogenic stem cells since they were CD44 (+).

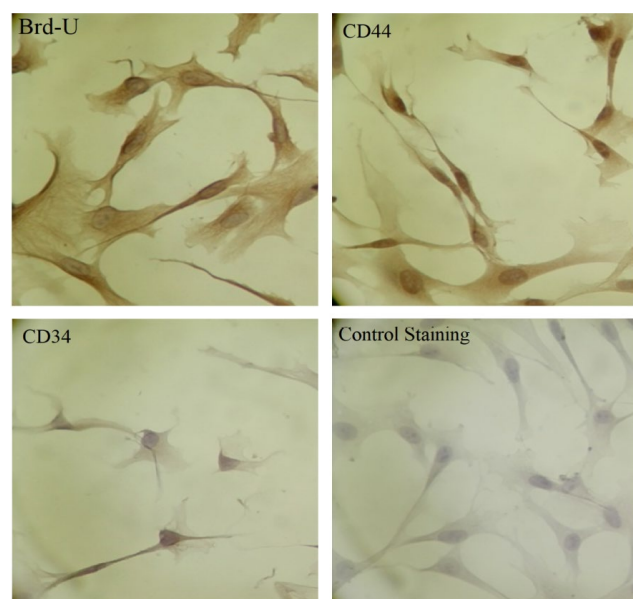


Figure 1. A: Brd-U, B: CD44, C: CD34, D: Control Staining

Histological Results

In the gross examination, the color of the ovaries in groups 1, 2, and 4 were less ischemic than group 3. Fibroblast accumulation around the ovarian follicles and small blood vessels filled with erythrocytes in the ovarian stroma were observed in group 1, whereas these changes decreased after ADSC administration in group 2. Intensely detected stromal fibroblast accumulation, infiltration and apoptotic cells in group 3 decreased after ADSC administration in group 4 than any other groups (Figure 2).

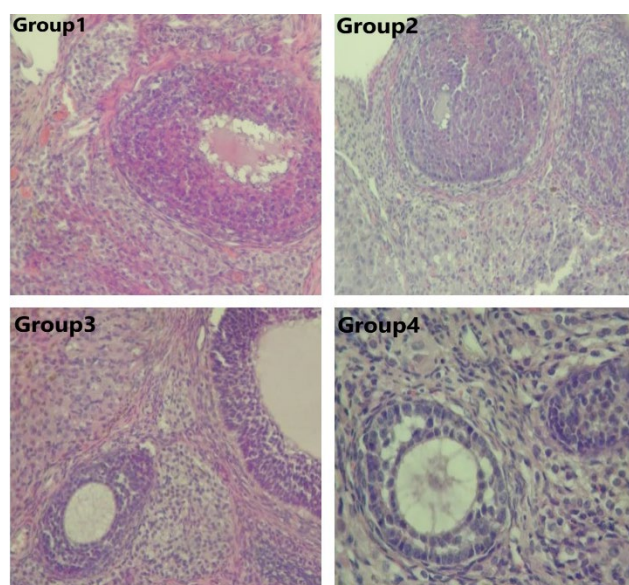


Figure 2. Histological analysis of ovarian tissue from group 1, Group 2, Group 3 and Group 4: X400 original magnification

Immunohistochemical Results

TUNEL analysis of ovarian tissue apoptosis is shown in Figure 3. No TUNEL-positive cells were observed in primary, secondary, or Graafian follicles in groups 1 and 2. However, apoptotic cells were detected in corona radiata and cumulus cells of Graafian follicles, and primary follicles of group 3. In group 4, TUNEL-positive cells were detected only in Graafian follicles, but in lower numbers than in group 3.

Immunohistochemical distributions of iNOS, eNOS, Caspase-3, and BrdU are shown in Figure 3, respectively. TUNEL, iNOS, eNOS, Caspase-3, and BrdU staining levels in the groups are also demonstrated in Table 1. BrdU staining was positive only in groups 2 and 4, confirming the presence of ADSCs in ovarian tissue after stromal injection.

Table 1. Staining levels of TUNEL, iNOS, eNOS, Caspase-3, and BrdU in experimental groups

Groups	TUNEL	iNOS	eNOS	Caspase-3	BrdU
Group 1	-	+ / ++	-	+	-
Group 2	-	++ / +++	-	++	+
Group 3	++	+++ / ++++	-	++	-
Group 4	+	++	-	+	+

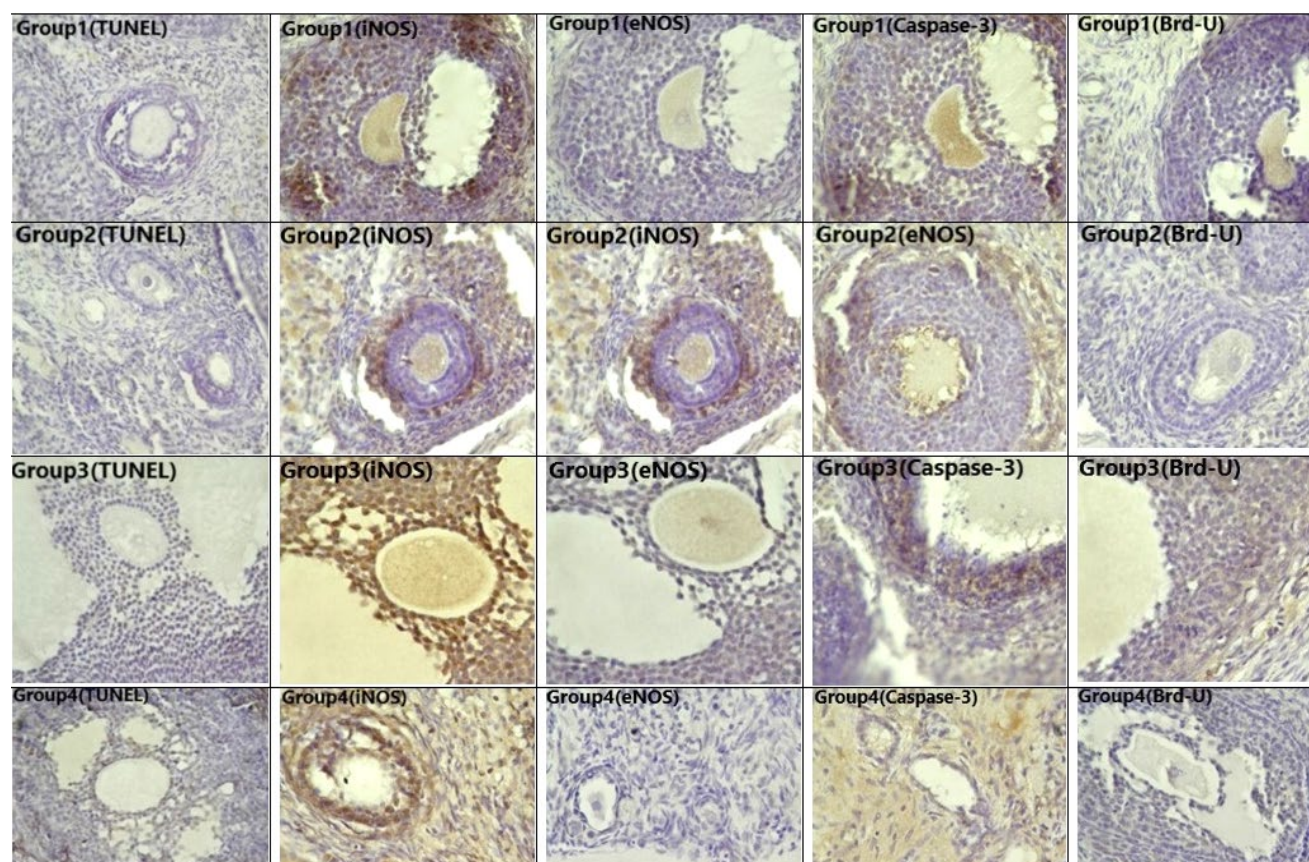


Figure 3. Apoptotic analysis of ovarian tissue from group 1, group 2, group 3 and group 4 after TUNEL assay: X400. Immunohistochemical distribution of iNOS, eNOS, caspase-3 and Brd-u. X400

Biochemical Results

AOPP levels in tissue and serum of all groups are presented in Tables 2 and 3, respectively. Levels of tissue AOPP were higher in group 3 than in group 2 ($p<0.05$), whereas levels of AOPP in group 4 were lower than those in groups 1 and 3 ($p<0.05$) (Table 2). Serum AOPP was higher in group 3 than in group 1 and 2 ($p<0.05$), whereas they significantly decreased compared to group 3 after treatment with

ADSCs in group 4 ($p<0.05$) (Table 3). SOD and MPO activities and TSH levels in tissue and serum in all groups are presented in Tables 2 and 3, respectively. Tissue SOD activity was significantly lower in group 3 than in group 1 ($p<0.05$). Additionally, we found that SOD activity in group 4 was significantly lower than that in groups 1 and 2 ($p<0.05$) (Table 2). The MPO activity in ovarian tissue in group 3 was greater than in group 2 ($p<0.05$) (Table 2).

Table 2. Tissue AOPP, MPO, SOD, and TSH levels in experimental groups

Variables	Group 1	Group 2	Group 3	Group 4
tAOPP ($\mu\text{mol}/\text{mg protein}$)	99.12 $\pm$ 16.41	51.70 $\pm$ 1.5*	91.70 $\pm$ 9.7 ψ	56.31 $\pm$ 3.97* ϕ
tMPO ($\text{units}/\text{g tissue}$)	0.225 $\pm$ 0.011	0.206 $\pm$ 0.027	0.393 $\pm$ 0.049 ψ	0.310 $\pm$ 0.049
tSOD ($\text{units}/\text{mg protein}$)	73.23 $\pm$ 10.02	58.14 $\pm$ 4.15	42.52 $\pm$ 7.29*	34.46 $\pm$ 2.77* ψ
tTSH ($\mu\text{mol}/\text{mg protein}$)	115.9 $\pm$ 20.86	107.6 $\pm$ 8.66	85.51 $\pm$ 8.65	164.10 $\pm$ 31.54

* $p<0.05$ compared to **Group 1**, ψ $p<0.05$ compared to **Group 2**, ϕ $p<0.05$ compared to **Group 3**. Results are given as mean $\pm$ standard error. Differences among experimental groups were determined by ANOVA and Tukey's post hoc tests

Table 3. Serum AOPP, SOD, and TSH levels in experimental groups

Variables	Group 1	Group 2	Group 3	Group 4
sAOPP ($\mu\text{mol/L}$)	93.02 $\pm$ 12.22	104.34 $\pm$ 9.34	199.84 $\pm$ 27.80 [*] , ^{ψ}	80.11 $\pm$ 7.31 ^{ϕ}
sSOD (<i>units/L</i>)	19.30 (16.88-20.28)	19.41 (17.48-20.61)	19.20 (16.55-21.49)	19.77 (18.46-27.42)
sTSH ($\mu\text{mol/L}$)	538.75 $\pm$ 47.6	502.62 $\pm$ 32.85	545.59 $\pm$ 75.42	390.98 $\pm$ 31.8

*p<0.05 compared to **Group 1**, ^{ψ} p<0.05 compared to **Group 2**, ^{ϕ} p<0.05 compared to **Group 3**. Results are given as mean $\pm$ standard error or median (minimum-maximum). Differences among experimental groups were determined by ANOVA and Tukey's post hoc tests. Kruskal-Wallis test was used for analyzing non-parametric data

Discussion

It was hypothesized that since ADSCs have antioxidant and anti-inflammatory properties, they might improve the harmful effects of I/R injury caused by ovarian torsion/detorsion surgery and the present study confirmed this by demonstrating that tissue and serum AOPP, as well as tissue SOD levels, were decreased after ADSC implementation. So, adjuvant treatment with ADSCs in an ovarian torsion model may mitigate ovarian damage by decreasing ROS in both tissue and serum, reducing the number of apoptotic cells and the inflammatory process.

Many opinions have been reported regarding the effects and mechanism of ADSC on human infertility and the reproductive system recently. In a study by Manavella et al, ADSC differentiates into human vessels and promotes the secretion of VEGF when transplanted together with human ovarian tissue to mouse peritoneum using a two-step ovarian tissue grafting procedure (28). They stated that ADSC may potentially improve ovarian tissue quality and lifespan. In another of Manavella et al, authors demonstrated higher rates of oxygenation and vascularization of ovarian tissue with ADSC in the early post-grafting period, which correlate with increased follicle survival and reduced apoptosis (29). Moreover, Sun et al found that ADSC sheets transplanted into excised uteri promoted regeneration of the endometrium cells, muscle cells and stimulated angiogenesis, and also resulted in better pregnancy outcomes (30). As can be

understood from these data, ADSCs appear to have positive effects on the human reproductive system and infertility.

The elevated levels of ROS could cause some damages to vital macromolecules including DNA, protein, and lipids and they play a significant role in the development of some pathological events such as I/R-induced ovarian injury (31). Such deleterious effects might be represented as a decrease in the number of follicles (pre-antral, and antral), as well as an increase in the number of atretic bodies and apoptosis of granulosa cells in follicles (10). The balance between ROS and antioxidants influences follicular growth and maturation because antioxidant supplements by reducing ROS and inhibiting lipid peroxidation can lead to preventing cell damage, enhancing follicular development, and increasing the fertilization rate and embryo development. The accumulation of ROS influences apoptosis. Shoorei et al found that hesperidin treatment led to an increase and a decrease in the mRNA levels of the Bcl-2 gene (anti-apoptotic) and Bax gene (pro-apoptotic), respectively. Hesperidin was also found to inhibit the generation of ROS. AOPPs have been widely used as reliable biomarkers for oxidant-mediated protein damage due to their stability (33). Previously, elevated plasma or serum AOPP has been reported to be associated with deterioration in atherosclerosis (34). In this study, AOPP was measured in both serum and tissue to estimate oxidant-mediated protein

damage and conclusively it was found that ADSC treatment decreased the AOPP levels in ovarian tissue and serum following I/R injury.

SOD and sulfhydryl groups are important antioxidants that protect the cell against free radical toxicity (35). Consistent with previous studies, decreased tissue SOD activity was found in the ovarian I/R model (35). Shojafar et al. performed bilateral oophorectomy/autotransplantation with and without ADSC implementation into autografted ovaries in mice to create an experimental model of I/R injury (36). They also presented the decrease of SOD in autografted ovaries and the increase of SOD in the group of autograft+ADSC. However, SOD activity was not significantly affected by treatment with ADSCs in current study. Besides, tissue TSH levels were decreased in the groups that underwent ovarian torsion compared to the control group 1, and TSH levels were higher in group 2 than in other groups. Nonetheless, these findings were not statistically significant. Additionally, there were no significant differences in systemic SOD and TSH levels among all the groups.

During I/R, increased ROS and other biological stimulants activate inflammatory cells, leading to neutrophil influx into the affected site. MPO, released by activated neutrophils, causes the formation of ROS and HCIO (37). MPO activity was found to be significantly increased after ovarian I/R, but significant decrease in MPO activity after ADSC treatment was not observed. Histological analysis was consistent with this observation; lymphocyte infiltration gathered in group 3, and somewhat diminished following ADSC treatment.

Nitric oxide (NO) plays an essential role in vasodilation, immunity, inflammation and apoptosis. As a Janus molecule, NO employs both beneficial and deleterious effects depending on its isoforms, such as eNOS or iNOS (38). In this study, they found that ADSC treatment reduced iNOS expression in group 4

compared with 3. Several studies demonstrated that along with inflammation and apoptosis. Former researches revealed that the number of apoptotic cells was significantly increased in experimental models of ovarian I/R (8). In this study, apoptosis was evaluated immunohistochemically with TUNEL and caspase-3 staining. Manavella et al. grafted frozen-thawed human ovarian tissue of five different patients in severe combined immunodeficient mice and intraperitoneally transplanted ADSC loaded in fibrin. They reported that significantly fewer apoptotic follicles were detected in the ADSC-loaded group in the TUNEL assay (29). In this study, a reduced number of apoptotic cells with ADSC treatment was shown by TUNEL assay.

There are some similar studies published on testicular I/R injury. In the study of Moghimian et al. conducted on rats with testicular I/R injury, the level of testosterone increased and MDA level decreased significantly in the 5 h testicular torsion group receiving treatment with vitamin C. The authors concluded that this may be a result of the antioxidant effect of vitamin C (39). In the study of Shokoohi et al. conducted on rats receiving hesperidin treatment, hesperidin increased the serum levels of GPx and SOD enzymes and decreased the heightened concentrations of MDA in testicular tissue. Moreover, the authors stated that hesperidin decreased Bax and increased Bcl-2 expression levels. They also demonstrated a markedly diminished rate of apoptosis in germ cells. The authors concluded that this may be a result of the antioxidant effect of hesperidin (40). In the study of Ameli et al. conducted on rats with testicular torsion, the levels of SOD, GPx, and testosterone were significantly decreased in the torsion/detorsion group while these increased in therapeutic groups receiving tadalafil and verapamil. Tadalafil and verapamil can protect the testis tissue damage and replace the testicular function by suppressing oxidative stress after testicular torsion (41). In the current study, the authors found that oxidative stress in

ovarian tissue also increased after ischemia. Moreover, tissue SOD activity and apoptotic cell number were found to be significantly decreased in group 4. These findings might be the result of the antioxidant properties of ADSC, similar to the studies mentioned above.

In addition to investigations demonstrating the therapeutic efficacy of stem cells, tumor-tropic adipose stem cells have been shown to support tumor vasculature formation, and enhance ovarian cancer progression (42). The tumorigenic potential of ADSCs in breast cancer was investigated by Eterno et al and reported that ADSCs were not tumorigenic but they might possess a tumorigenic potential in presence of c-Met-expression (43). Thus, precautions are advised for the use of stem cells in benign gynecological disorders.

The power of our study might have been increased by using longer ischemia durations and by detecting ovarian reserve via follicular count, anti-Müllerian hormone, inhibin B, estradiol, or fecundity. However, the presence of only the ADSC-given group is a strong side of our study in terms of seeing ADSC effects on the undamaged ovary.

In conclusion, if administered within 6 hours of torsion, ADSC treatment might protect against ovarian I/R damage by decreasing protein oxidation and iNOS expression, attenuating lymphocyte infiltration, and decreasing the incidence of apoptotic cells. According to the results of the present study, a promising aspect of ADSCs might have lightened for human studies and finally, ADSCs can be used as a potential adjuvant treatment agent in ovarian torsion cases.

Conflict of Interest

The authors declared no conflict of interest.

Author Contributions

BK: contributed to conceiving the study design, performing experimental study, writing and drafting the present article. ŞK contributed to

conceiving the study design and performing experiment. SV: carried out histopathological and immunohistochemical analyses and contributed to critical revision of the article. RK carried out histopathological and immunohistochemical analyses. OY and MÇ contributed to performing experimental study. ÜÖT carried out biochemical and statistical analyses and drafted the article. MNA: contributed to writing and drafting the article. EA contributed to writing and drafting the article. NÖT: contributed to critical revision of the article.

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