Endometrial scratching for poor responders based on the Bologna criteria in ICSI fresh embryo transfer cycles: a preliminary retrospective cohort study

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Abstract

Objective: This study evaluated the effect of endometrial injury on pregnancy outcomes in patients with a poor ovarian response (POR), based on the Bologna criteria, who underwent intracytoplasmic sperm injection (ICSI) cycles.

Material and Methods: Sixty-eight patients were enrolled in this retrospective cohort study. All patients in the endometrial scratching group (group 1, n=32) and control group (group 2, n=36) underwent office hysteroscopy in the early follicular phase of the cycle before controlled ovarian stimulation. Group 1 also underwent endometrial scratching. The main outcome measure was the ongoing pregnancy rate.

Results: The study groups had similar baseline demographics, including age, body mass index, duration of infertility, number of ICSI cycles, and hormone levels. However, the antral follicle count was significantly higher in group 1 than in group 2 (4.2±1.9 vs 3.3±1.8; p<0.05). There were no significant group differences in ovarian stimulation characteristics (ovarian stimulation time, trigger day endometrial thickness, number of metaphase II oocytes), number of embryos transferred, or the ratio of embryo transfer on days 3 and 5. Moreover, there were no significant differences between groups 1 and 2 in the rates of chemical pregnancy (25% vs 19.4%), clinical pregnancy (15.6% vs 11.1%) or ongoing pregnancy (9.4% vs 8.3%) (p>0.05 for all).

Conclusion: Endometrial scratching did not improve pregnancy outcomes for patients meeting the Bologna criteria for a POR to ICSI cycles using fresh embryo transfer and the GnRH antagonist protocol. (J Turk Ger Gynecol Assoc 2021; 22: 47-52)

Keywords: Endometrial scratching, Bologna criteria, poor ovarian response, ICSI, GnRH antagonist protocol

Introduction

The world's first in-vitro fertilization (IVF) baby was born in 1978. Since then, according to the International Committee for Monitoring Artificial Reproductive Technology, approximately 8 million babies have been born worldwide by means of IVF or other advanced infertility treatment methods. However, pregnancy rates in Europe appear to have stabilized, at approximately 36%, for both IVF and intracytoplasmic sperm injection (ICSI) treatment cycles (1).

The number of oocytes retrieved during an IVF cycle (ideally about 15) is directly related to the success thereof (2). A poor ovarian response (POR), in which controlled ovarian stimulation (COS) during IVF/ICSI cycles yields a limited number of oocytes, has an average prevalence of 6% to 35% (3,4). POR is frustrating for both patients and clinicians, as it is related to low pregnancy rates in IVF and high cancellation rates during COS (5). Thus, strategies to increase pregnancy rates in patients with POR are important in IVF/ICSI cycles.
Embryo implantation is one of the most important steps in the IVF cycle. Implantation failure may be caused by low endometrial receptivity, embryo problems, or abnormalities of both the endometrium and embryo. As reported previously, about 66% of implantation failures are caused by decreased endometrial receptivity (6). Thus, ensuring a receptive endometrium could increase pregnancy and delivery rates for artificial reproductive technology (ART). Various techniques have been used to ensure a receptive endometrium, including management of intracavitary abnormalities by hysteroscopy, treatment of thin endometrium, immunotherapy, and adjuvant treatments for women with POR (7,8).

Endometrial scratching, which is also called endometrial biopsy, endometrial injury, and endometrial trauma, has been offered by clinicians as a means of increasing endometrial receptivity, by an as yet unknown mechanism (9). Endometrial scratching can be done with endometrial biopsy instruments during the early follicular or luteal phase of the preceding IVF treatment cycle. The first study demonstrating a substantial increase in pregnancy rates after endometrial scratching was followed by many additional studies and reviews including various populations, although these have had conflicting results (10-15). However, no study has directly addressed the effect of endometrial scratching on patients with POR. The objective of this preliminary trial was to assess the role of endometrial scratching in patients with a POR undergoing ICSI cycles with fresh embryo transfer.

**Material and Methods**

This study included 68 women seen at a private IVF Center in Koru Hospital, Ankara, Turkey. Approval was obtained from the Koru Hospital Ethical Committee before the study commenced (approval number: 81, date: 04/11/2019). The study population was enrolled between January 1, 2017 and November 1, 2019. The Declaration of Helsinki was followed.

All participants were undergoing their first or repeated ICSI fresh embryo transfer and met the following inclusion criteria: fulfillment of the Bologna criteria for POR (16); aged 20-42 years; body mass index (BMI) 20-30 kg/m²; and a normal uterine cavity based on office hysteroscopy or hysterosalpingography. Patients were excluded if they had uterine anomalies, endocrine disorders, ovarian cysts, hydrosalpinx, or severe male factor infertility (e.g. aspermia, azoospermia), or if they had undergone uterine surgery in the last 3 months, or IVF cycles for preimplantation genetic diagnosis. Patients with genetic abnormalities, and those who had undergone cryo-thawed embryo transfer, were also excluded.

A detailed history, including age, previous treatments, and duration of infertility, was taken from all patients. A gynecologic examination consisting of a bimanual pelvic examination plus transvaginal ultrasonography (TVUSG) was conducted to check for structural abnormalities of the pelvis and ovaries. On days 2-5 of the menstrual cycle, gonadotropic hormone and estradiol concentrations were assessed.

All patients underwent office hysteroscopy during the early follicular phase of the menstrual cycle, i.e., immediately preceding the planned IVF cycle. A rigid 30°, 5.5 mm hysteroscope was used to perform hysteroscopy without anesthesia (Karl Storz Endoscopy, Tuttingen, Germany). Serum physiological solution was used to distend the uterine cavity during hysteroscopy. Group 1, but not group 2, also underwent endometrial scratching during hysteroscopy; scissors were used to create mechanical tissue damage in local areas of the fundus, and in posterior and anterior regions of the endometrium.

COS began on day 2 or 3 of the menstrual cycle. Subcutaneous injections of recombinant follicle-stimulating hormone (FSH) (Gonal-F; Serono, Rome, Italy) or highly purified human menopausal gonadotropin (Menopur, Ferring, Sweden or Merional; IBSA, Collina d’Oro, Switzerland) were used for ovarian stimulation. The initial dose for each patient was based on the predicted ovarian response, and varied from 300 to 450 IU. On day 5 of COS, the flexible GnRH antagonist treatment protocol was implemented to prevent premature luteinizing hormone (LH) surge (Cetrotide; Merck Sharp and Dohme Ltd., Athens, Greece). Hormones, including FSH, estradiol, and thyroid-stimulating hormone, were measured before stimulation. LH, estradiol, and progesterone were also measured on the day of human chorionic gonadotropin (hCG) administration.

When two or more follicles with a diameter of at least 17 mm were detected, ovulation was triggered by a subcutaneous injection of 250 µg r-hCG (Ovitreel; Serono). Oocyte retrieval was carried out 35.5-36 hours after the r-hCG injection using TVUSG. On day 3 or day 5 after oocyte retrieval, depending on oocyte development, a maximum of two good-quality embryos were transferred into the uterine cavity using a Wallace semirigid catheter (Cooper Surgical, Malov, Denmark) under abdominal ultrasonography guidance.

Luteal support was provided until the 12th week of gestation using Crinone gel (8% progesterone Serono). A pregnancy test was conducted approximately 2 weeks after embryo transfer. Clinical pregnancy was defined as any intrauterine gestational sac with a fetal heartbeat at 4 weeks after the first pregnancy test. The existence of at least one live fetus at the 12th week of gestation was considered an ongoing pregnancy, which was the primary outcome measure.

**Statistical analysis**

Statistical analysis was carried out using SPSS software (version 23.0; SPSS Inc., Chicago, IL, USA). Continuous
variables are expressed as mean ± standard deviation or median (minimum-maximum). Categorical variables are expressed as number and percentage (%). The Kolmogorov-Smirnov test was used to check the distribution of the data. The Independent Samples t-test and Mann-Whitney U test were used to compare continuous variables. Pearson’s chi-squared test or Fisher’s exact test was used to compare categorical variables. A two-tailed p-value of <0.05 was considered significant.

Results

A total of 68 patients were included in the present study. The endometrial scratching group (group 1) and control group (group 2) included 32 and 36 women, respectively. The groups were well-balanced in terms of baseline demographic data, including age, BMI, length of infertility, number of IVF cycles, and hormone levels (Table 1). However, the antral follicle count was significantly higher in group 1 compared to group 2 (4.2±1.9 vs 3.3±1.8; p<0.05).

Table 2 shows the COS, ICSI, and pregnancy outcomes. Groups 1 and 2 did not differ in duration of COS, trigger day endometrial thickness, or number of metaphase II oocytes retrieved. In addition, the number of embryos transferred, and the ratio of day 3 to day 5 embryo transfers, were similar between the two groups. Moreover, regarding pregnancy outcomes, groups 1 and 2 had similar rates of chemical pregnancy (25% and 19.4%, respectively), clinical pregnancy (15.6% and 11.1%) and ongoing pregnancy (9.4% and 8.3%).

Discussion

This retrospective cohort trial investigated the impact of endometrial scratching on the pregnancy outcomes of women meeting the Bologna criteria for POR, who were undergoing ICSI fresh embryo transfer cycles using the GnRH antagonist protocol. The results showed that endometrial scratching did not increase rates of clinical or ongoing pregnancy.

Poor responders constitute a major challenge for ART. These patients are more likely to show poor oocyte retrieval and less favorable pregnancy outcomes. A previous review reported that 47 studies used 41 different definitions of POR (17). To overcome this challenge, ESHRE used the Bologna criteria to clearly define POR (16).

Endometrial injury may be useful in patients undergoing ART cycles, to improve endometrial receptivity and the chance of pregnancy. Although endometrial injury has been studied for over a decade, the biological mechanism by which it increases the chance of pregnancy is not clear. However, one putative mechanism is stimulation of the production of cytokines and growth factors, which are essential for endometrial receptivity and embryo implantation after endometrial scratching and subsequent healing (18).

Endometrial injury to improve the outcomes of IVF/ICSI cycles has attracted increasing attention since Granot et al. (19) conducted the first two studies in 2000 and 2003 (20). In the first study, repeated endometrial biopsies were performed to evaluate the endometrium of 12 infertile women with several unsuccessful cycles of IVF treatment. The authors noted that 11 of these women became pregnant during the first IVF cycle after endometrial biopsy (19). In the second study, a quasi-

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Table 1. Baseline demographic, clinical, and laboratory characteristics of groups 1 and 2

| Parameter                   | Group 1 (n=32) | Group 2 (n=36) | p     |
|-----------------------------|----------------|----------------|-------|
| Age (years)                 | 34.1±4.0       | 33.2±4.0       | 0.361 |
| Body mass index (kg/m²)     | 24.7±3.4       | 25.4±3.8       | 0.754 |
| Duration of infertility (years) | 7.2±3.8       | 5.3±2.4       | 0.045 |
| Number of cycles (n)        | 1.6±0.8        | 1.6±0.8        | 0.854 |
| Antral follicle count (n)   | 4.2±1.9        | 3.3±1.8        | 0.035 |
| Baseline FSH (IU/L)         | 10.2±3.0       | 11.8±3.7       | 0.123 |
| Baseline estradiol (pg/mL)  | 50.5±19.0      | 52.1±19.4      | 0.736 |

Data are presented as mean ± standard deviation, median (interquartile range), or number (percentage). FSH: Follicle stimulating hormone, a: Mann-Whitney U test, b: Independent samples t-test, *Statistically significant difference
Table 2. Controlled ovarian stimulation and pregnancy outcomes of groups 1 and 2

| Parameter                      | Group 1 (n=32) | Group 2 (n=36) | p    |
|--------------------------------|---------------|---------------|------|
| Stimulation period (days)      | 8.3±1.5       | 7.8±1.5       | 0.146a|
|                               | 8.0 (7.0; 9.0) | 8.0 (7.0; 8.0) |      |
| Trigger day endometrial thickness (mm) | 8.8±1.7   | 9.1±1.9       | 0.397b|
|                               | 8.7 (7.2; 9.9) | 8.8 (7.6; 10.0) |      |
| MI oocytes (n)                 | 2.4±1.3       | 2.6±1.4       | 0.622a|
|                               | 2.0 (1.0; 3.0) | 2.0 (1.5; 4.0) |      |
| 2PN embryos (n)                | 1.8±1.0       | 1.9±1.0       | 0.486a|
|                               | 1.5 (1.0; 2.0) | 2.0 (1.0; 2.0) |      |
| Embryos transferred (n)        | 1.3±0.5       | 1.3±0.4       | 0.772a|
|                               | 1.0 (1.0; 2.0) | 1.0 (1.0; 1.5) |      |
| Embryo transfer day, n (%)     |               |               | 0.725c|
| 3                              | 27 (84.4%)    | 32 (88.9%)    |      |
| 5                              | 5 (15.6%)     | 4 (11.1%)     |      |
| Pregnancy outcomes, n (%)      |               |               |      |
| Beta hCG positive              | 8 (25%)       | 7 (19.4%)     | 0.581d|
| Clinical pregnancy             | 5 (15.6%)     | 4 (11.1)      | 0.725c|
| Ongoing pregnancy              | 3 (9.4%)      | 3 (8.3%)      | 1.000c|

Data are presented as mean ± standard deviation, median (interquartile range) or number (percentage).

hCG: Human chorionic gonadotropin, PN: Pronucleus, M: Metaphase, a: Mann-Whitney U test, b: Independent Samples t-test, c: Fisher’s exact test, d: Pearson’s chi-squared test.
scratching techniques and COS protocols, and larger samples drawn from this subgroup of patients, are needed.

**Ethics Committee Approval:** Approval was obtained from the Kura Hospital Ethical Committee before the study commenced (approval number: 81, date: 04/11/2019).

**Informed Consent:** Retrospective study.

**Peer-review:** Externally and internally peer-reviewed.

**Author Contributions:** Surgical and Medical Practices: Ş.K., A.Y., M.D.; Concept: Ş.K., A.Y., M.D.; Design: Ş.K., A.Y., M.D.; Data Collection or Processing: Ş.K., A.Y., M.D.; Analysis or Interpretation: Ş.K., A.Y., M.D.; Literature Search: Ş.K., A.Y., M.D.; Writing: Ş.K., A.Y., M.D.

**Conflict of Interest:** No conflict of interest is declared by the authors.

**Financial Disclosure:** The authors declared that this study received no financial support.

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