https://doi.org/10.5653/cerm.2021.04497 plSSN 2233-8233 • elSSN 2233-8241 Clin Exp Reprod Med 2021;48(4):362-367



Comparison of embryonic competence and clinical outcomes between early and late cumulus cell removal for *in vitro* fertilization

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Objective: The impact of early mechanical removal of cumulus cells on fertilization and embryonic development is not yet precisely known. This study aimed to investigate the effects of early and late cumulus cell removal on fertilization, polyspermy, embryonic development potential, blastocyst development, and clinical outcomes.

Methods: A prospective study was conducted of patients who underwent *in vitro* fertilization between September 2019 and October 2020. Sibling oocytes were randomly allocated after insemination to early cumulus cell removal at 6 hours (group I) and late cumulus cell removal at 16–18 hours (group II). If total fertilization failure (TFF) was determined to have occurred at early cumulus cell removal, rescue intracytoplasmic sperm injection (ICSI) was performed. Fertilization, embryonic development, and pregnancy outcomes were compared.

Results: A total of 912 oocytes were assigned to group I (458 oocytes) and group II (454 oocytes). Fertilization, polyspermy, embryo quality, and pregnancy outcomes were not significantly different between both groups. Rescue ICSI enabled fertilization of 79.2% of the TFF oocytes. **Conclusion:** Early cumulus cell removal at 6 hours had no significant difference in fertilization, polyspermy, embryo development, or obstetric and perinatal outcomes compared to late removal. Early cumulus cell removal combined with early rescue ICSI may have the potential to help couples with TFF.

Keywords: Cumulus cell; In vitro fertilization; Embryo development

Introduction

Cumulus cells play a significant role in supporting oocytes and fertilization. In natural conception, cumulus-enclosed oocyte and spermatozoa meet together at the ampulla of the uterine tube. Cumulus cells enhance capacitation, sperm binding and penetration of the zona pellucida, passage through the perivitelline space, binding and fusion to the oolemma, activation of the cortical reaction, and formation of the male pronucleus. They can also provide nutrients, hor-

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The impact of early mechanical denudation of cumulus cells on fertilization and embryonic development is not yet precisely known. Early cumulus cell removal has been claimed to yield better embryo quality [4,5], but some studies showed comparable results [6,7] and

Received: March 15, 2021 · Revised: May 25, 2021 · Accepted: May 31, 2021 Corresponding author: **Pallop Pongsuthirak**

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others revealed conflicting outcomes [8]. However, an additional benefit of early cumulus cell removal is early recognition of fertilization based on the presence of a second polar body, and rescue ICSI can be conducted in cases of failed or low fertilization with promising outcomes [7-9]. However, there is still no consensus regarding early cumulus cell removal. This study aimed to investigate the effects of early cumulus cell removal compared to late cumulus cell removal on fertilization, polyspermy, embryonic development potential, blastocyst development, and clinical outcomes.

Methods

1. Study design and patient selection

This prospective randomized clinical study was carried out at Buddhachinaraj Hospital between September 2019 and October 2020. The study was registered with the Thai Clinical Trial Registry (TCTR 20190817001) and approved by the Research Ethics Committee of Buddhachinaraj Hospital Medical School (IRB 066/62). Patients who attended the infertility clinic and had indications for assisted reproduction were invited to join the study and provided written informed consent after enrollment. The inclusion criteria were women undergoing their first IVF treatment cycle aged 20-38 years old, who had at least 6 retrieved oocytes and whose partners had normal semen parameters. The etiologies of infertility included ovulatory dysfunction, tubal disease, unexplained infertility, and endometriosis. The oocytes were randomly allocated into two groups: early cumulus cell removal (group I), in which cumulus cells were removed 6 hours after insemination, and late cumulus cell removal (group II), in which cumulus cells were removed 16-18 hours after insemination. Rescue ICSI was conducted when total fertilization failure (TFF) was expected to occur in the early cumulus cell removal group 6 hours after insemination. Embryo transfers were selected randomly in the two groups in terms of whether the transferred embryos originated from the early or late cumulus cell removal group.

2. Controlled ovarian hyperstimulation and oocyte retrieval

Controlled ovarian hyperstimulation was started with a gonadotropin-releasing hormone agonist for down regulation in the mid-luteal phase of the previous cycle. Follicle-stimulating hormone and/or human menopausal gonadotropin in individually adjusted doses were administered after pituitary desensitization. If at least three follicles were \geq 18 mm, 5,000– 10,000 IU of human chorionic gonadotropin was injected to induce ovulation. The oocytes were then retrieved with a single-lumen needle under transvaginal ultrasound guidance 36–38 hours later.

3. Sperm preparation and insemination

Semen samples were gathered by masturbation in the morning on the same day of oocyte collection following 3–5 days of sexual abstinence. Sperm concentration, motility, and morphology were examined under a light microscope based on the World Health Organization criteria (fifth edition, 2010). Gradient centrifugation was used for sperm preparation. Active motile spermatozoa were harvested for insemination, with 20,000–50,000 spermatozoa contained in 50 μ L of insemination medium.

4. Cumulus cell removal

The oocytes were transferred to a new sperm-free medium after 4 hours of co-incubation. In group I, the cumulus cells were mechanically removed at 6 hours post-insemination using a denuding pipette (Flexipet; Cook, Brisbane, Australia) with an inner diameter of 140 µm under an inverted microscope. Fertilization was considered to have occurred when a second polar body was present, and TTF was deemed to have occurred when in the absence of a second polar body in any of the mature oocytes. Rescue ICSI was done if none of the oocytes showed early fertilization after insemination for 6 hours. In group II, the cumulus cells were removed at 16–18 hours after insemination.

5. Fertilization assessment and embryo culture

The developmental competence of zygotes was evaluated after 96–120 hours. Normal fertilization was defined as the presence of 2 pronuclei and polyspermy as the presence of \geq 3 pronuclei. Embryos were placed in cleavage medium during days 1–3 after fertilization, followed by blastocyst medium during days 4–5. Embryo morphology was assessed on day 5. The blastocysts were assigned a score based on the Gardner system, with high-quality blastocysts having scores of \geq 4 BB [10]. The surplus high-scoring blastocysts were cryopreserved for future transfers.

6. Clinical and birth outcomes

Embryo transfer took place on day 5 under ultrasound guidance. The number of embryos was limited to one blastocyst to reduce the risk of multiple pregnancies. Progesterone was started on day 3 after oocyte retrieval for luteal support. The implantation rate was defined as the number of gestational sacs divided by the number of embryos that were transferred. The fetal heartbeat demonstrated by ultrasonography was considered to be clinical pregnancy at 5 weeks after embryo transfer. Information was gathered on ongoing pregnancy at 20 weeks and the birth outcomes. Premature delivery was defined as a baby born before 37 weeks.

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7. Statistical analyses

Continuous data were compared using the Student *t*-test, and proportional data were compared with the chi-square test and the Fisher exact test. A *p*-value < 0.05 was considered to indicate statistical significance. The analyses were performed using SPSS ver. 13.0 (SPSS Inc., Chicago, IL, USA). A sample size calculation found that at least 240 oocytes in each arm would be satisfactory for a 10% difference in maturation rates between oocytes in the two groups, given a type I error of 5% (two-tailed) and a type II error of 20%.

Results

A total of 85 patients were eligible for the study. Seven patients were excluded due to TTF, and rescue ICSI was performed (Figure 1). The remaining 78 patients were 32.8 ± 2.6 years old. Thirty-three patients (42.3%) had tubal factor fertility, 8 (10.2%) had endometriosis, 10 (12.8%) had polycystic ovary syndrome, and 27 (34.7%) had unexplained infertility (Table 1). In total, 912 oocytes were randomly allocated to early cumulus cell removal (group I, 458 oocytes) and late cumulus cell removal (group II, 454 oocytes). The details of fertiliza-



Figure 1. Consolidated standards of reporting trials (CONSORT) flowchart. ICSI, intracytoplasmic sperm injection.

tion and embryo development are summarized in Table 2. No significant differences were observed in fertilization, cleavage rate, and embryo quality. Polyspermy was higher in group I than in group II (9.0% vs. 5.9%, respectively), but the difference did not reach statistical significance.

Embryo transfer was conducted in 78 patients, of whom 40 patients obtained embryos from group I and 38 patients from group II. There were no significant differences between the two groups in terms of age, and the number of oocytes. Implantation rates, clinical pregnancy, ongoing pregnancy, and live birth rate were not significantly different between the patients with embryos transferred from the early and late cumulus cell removal groups (Table 3). No twin pregnancies occurred in this study.

Notably, seven patients in the early cumulus cell removal group had TTF, and rescue ICSI was carried out. Nineteen of 24 oocytes (79.2 %) had normal fertilization without detected polyspermy and three embryos were transferred in three cycles leading to one single pregnancy with an uneventful course of gestation and delivery of a normal newborn (Table 4). However, in the late cumulus cell removal group, four patients had simultaneous fertilization in the sibling oocytes of the same patients, the embryos were transferred with a suc-

Table 1. Baseline characteristics of the 78 patients who participated in the study

Characteristics	Value
Age (yr)	30.4 ± 3.7
Body mass index (kg/m²)	25.5 ± 2.6
Etiology of infertility	
Tubal factor	33 (42.3)
Endometriosis	8 (10.2)
PCOS	10 (12.8)
Unexplained infertility	27 (34.7)

Values are presented as mean±standard deviation or number (%). PCOS, polycystic ovarian syndrome.

Table 2. Outcomes of oocyte and embryo development using early	y
and late cumulus cell removal	

Variable	Early removal (group I)	Late removal (group II)	<i>p</i> -value ^{a)}
Mature oocyte	458	454	-
Two pronuclei	375 (81.9)	373 (82.2)	0.913
Polyspermy	41 (9.0)	27 (5.9)	0.111
Cleavage	320 (85.3)	302 (81.0)	0.112
Blastocyst formation	178 (55.6)	156 (51.7)	0.323
High quality blastocyst	87 (27.1)	71 (23.5)	0.294

Values are presented as number (%).

^{a)}Chi-square test.



Table 3. Pregnancy outcomes according to early and late cumulus cell removal

Variable	Early removal (group I)	Late removal (group II)	<i>p</i> -value
No. of patients	40	38	-
Mean age (yr)	32.2±2.7	33.3±2.5	0.066 ^{a)}
No. of embryos transferred	1	1	-
Implantation rate	19/40 (47.5)	19/38 (50.0)	0.995 ^{b)}
Clinical pregnancy rate	18/40 (45.0)	17/38 (44.7)	0.838 ^{b)}
Ongoing pregnancy rate	16/40 (40.0)	15/38 (39.5)	0.854 ^{b)}
Live birth rate	14/40 (35.0)	14/38 (36.8)	0.947 ^{b)}
Premature delivery rate	2/16 (12.5)	1/15 (6.7)	0.999 ^{c)}
Birth weight (g)	$2,930 \pm 266$	3,040±250	0.059 ^{a)}

Values are presented as mean±standard deviation or number (%).

^{a)}Independent *t*-test; ^{b)}Chi-square test; ^{c)}Fisher exact test.

Table 4. Details of embr	ryonic competence and	l clinical outcomes in seven	patients who had rescue ICSI
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Patient	Age (yr)	Oocyte	Fertilization	Cleavage	Blastocyst	Implantation	Clinical pregnancy	Birth
1 ^{a)}	28	4	3	3	2	0	0	0
2 ^{a)}	30	3	3	3	3	1	1	1
3 ^{b)}	31	4	4	3	1	-	-	-
4 ^{b)}	31	3	2	2	2	-	-	-
5 ^{b)}	32	3	2	2	1	-	-	-
6 ^{a)}	34	4	3	2	2	0	0	0
7 ^{b)}	37	3	2	2	1	-	-	-

ICSI, intracytoplasmic sperm injection.

^{a)}Patients 1,2, and 6 had a rescue ICSI-derived embryo transferred. Patient 2 became pregnant and delivered a normal newborn; ^{b)}Patients 3,4,5, and 7 did not have a rescue ICSI-derived embryo transferred.

Table 5. Details of embryonic competence and clinical outcomes of late cumulus removal oocytes in four patients who had TFF of sibling oocytes of early cumulus removal

Patient	Age (yr)	Oocyte	Fertilization	Cleavage	Blastocyst	Implantation	Clinical pregnancy	Birth
3	31	4	3	3	2	0	0	0
4 ^{a)}	31	4	3	3	3	1	1	1
5 ^{b)}	32	4	4	2	2	0	0	0
7	37	3	2	2	1	0	0	0

TFF, total fertilization failure.

^{a)}Patient 4 had a pregnancy with breech presentation, and the infant was delivered by cesarean section; ^{b)}Patient 5 had two polyspermic-fertilized zygotes

cessful singleton, and there was a normal course of pregnancy with breech presentation, followed by delivery of a normal infant by cesarean section (Table 5).

Discussion

There are inconsistent reports on the outcomes of early cumulus cell removal. Some studies have shown that early cumulus cell removal has fertilization and clinical pregnancy rates similar to late removal at 20 hours [6,7]. Another study demonstrated that early cumulus removal after 4 hours of insemination resulted in low numbers

of available embryos [8]. However, a meta-analysis [11] showed that early cumulus cell removal was associated with a significant increase in the implantation rate and clinical pregnancy rate. The ability to draw a clear conclusion may be hindered by differences in study designs and populations. In the present study, sibling oocytes were randomly allocated into two groups to minimize the confounding between patients, and the findings indicated that early cumulus cell removal had no significant difference in fertilization, polyspermy, cleavage of embryos, clinical pregnancy, and ongoing pregnancy, comparable with the findings of previous studies [4,5,12-16].

Conventionally, cumulus cells are recognized as essential for oo-



cyte development and the natural fertilization process, but become less important after ICSI because embryos can develop normally without cumulus cells. This fact implies that the early removal of cumulus cells may not affect embryonic development. Moreover, Nagy et al. [17] found that oocvtes were fertilized 2-4 hours after exposure to spermatozoa and the second polar body was extruded by approximately 90% into perivitelline space by 6 hours [6]. If this event is observed during early cumulus cell removal, rescue ICSI can be performed, resulting in higher fertilization rates and optimal embryos compared with rescue ICSI after late cumulus cell removal (20 hours) [6]. Interestingly, the time-course of fertilization in early rescue ICSI has a similar pattern to those oocytes that undergo ICSI at the normal time of fertilization, allowing the embryos to be obtained in synchronized development with the endometrium [6]. In this study, cumulus cells were dissected at 6 hours and rescue ICSI was conducted in seven cases of TFF, which obtained a satisfactory fertilization rate (79.2%); thus, meant ICSI after early cumulus cell removal (6 hours) could rescue most of the unfertilized oocytes. Therefore, early cumulus cell removal in conjunction with rescue ICSI provides an additional benefit by alleviating cycle cancellation in patients with TFF. Nevertheless, it is possible that injecting fertilized eggs can delay extrusion of the second polar body (3.2%) [9]. To avoid such an event, rescue ICSI can be postponed to 9 hours after insemination [12]. Remarkably, polyspermy after rescue ICSI was not found even when performed at 6 hours in our study. Despite the potential advantage of early cumulus cell removal and rescue ICSI, the demand of embryologists to cover the additional work during the extra period must be considered [6,11].

This study found a trend toward a higher polyspermy rate, albeit without statistical significance, in early cumulus cell removal (9.0%) than in late removal (5.9%), which aligns with previous reports [7,15]. It is unclear whether early cumulus cell removal can affect polyspermy. Cumulus cells are tougher and more difficult to remove at the earlier time point than at the later time point. However, at an early period after insemination, the oocytes are also more vulnerable due to their active spindles and microtubules [18,19]. Thus, they may have more susceptible to damage from the additional mechanical force created by the denuding pipette during cumulus cell removal [16]. Additionally, repeated mechanical stress can also have adverse effects on the integrity of the zona pellucida and may reduce the protective mechanism against polyspermic fertilization [16]. Of particular note, unstable culture conditions or an excessive number of sperm in the culture medium can also affect abnormal fertilization through different pathways [12]. These possible mechanisms may contribute to an increase in polyspermy. However, a meta-analysis showed that the use of a denuding pipette during cumulus cell removal was not harmful to the clinical pregnancy or implantation rate [11].

Our results showed that early cumulus cell removal had comparable obstetric and prenatal outcomes with late cumulus cell removal, in agreement with Liu et al. [14]. However, recent studies [11,13] reported higher rates of premature delivery, twins, and low-birthweight newborns in patients who underwent early cumulus cell removal. They proposed that the process during removal of cumulus cell at the early time point could possibly alter spindle integrity and impair cell division more than late removal, which might give rise to twins or poor fetal growth [20]. In addition, inappropriate mechanical forces during cumulus cell removal can cause epigenetic changes associated with low birth weight [21,22]. Further studies should be conducted to elucidate this interesting issue.

The present study showed that early cumulus cell removal at 6 hours after insemination had no significant difference in fertilization, polyspermy, embryo development, obstetric and perinatal outcomes. Early cumulus cell removal combined with early rescue ICSI may have the potential to help couples with TTF.

Conflict of interest

No potential conflict of interest relevant to this article was reported.

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