

Original Article

Comparison of Unstimulated In Vitro Maturation and Stimulated In Vitro Fertilization in Women with Poly Cystic Ovarian Syndrome

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ABSTRACT

Background: In regard to complications of controlled ovarian hyperstimulation (COH) such as multiple pregnancy and ovarian hyper stimulation syndrome (OHSS) and other short and long term complications, the interest to In Vitro Maturation (IVM) of oocytes from unstimulated ovaries is increasing. This clinical trial study was designed to establish the relative success of treatment with IVM in comparison with In Vitro Fertilization (IVF) in Poly Cystic Ovarian Syndrome (PCOS) women.

Methods: This prospective clinical trial included 35 IVM and 35 IVF cycles matched for age, and duration and cause of infertility. IVM patients underwent transvaginal retrieve of immature oocytes during unstimulated cycles, IVM and ICSI. Those in IVF group underwent ovarian stimulation after pituitary suppression. Main outcome measures included number of retrieved oocytes, number of produced embryos, and pregnancy rate and complications.

Results: In the IVM group, 1.39 mature oocytes, 0.27 fertilized oocyte, and 0.24 embryos were obtained per retrieval, and in IVF group, 8.45 mature oocytes, 3.8 fertilized oocytes, and 2.7 embryos were obtained.

Conclusion: This significant differences between two groups ($P < 0$) indicate that this method maybe used as a research setting until further study.

Key words: In Vitro Maturation, ART, unstimulated IVM, PCOS.

In vitro fertilization (IVF) is associated with two major complications: multiple pregnancy and OHSS due to controlled ovarian hyperstimulation (COH) especially in PCOS women. COH isn't required for In vitro maturation (IVM), Therefore, this method is cheaper and safer than IVF. In addition, regard to short and long term complications and side effects of COH, such as abdominal pain, nausea, and vomiting, unknown effects on dysplastic changes of ovaries³⁷, IVM of oocytes from unstimulated ovaries is a reproductive technology of increasing interest³⁵. Indications for IVM include: PCOS women requiring IVF, primarily poor quality embryos in repeated previous IVF without apparent reason, poor responders to high dose gonadotrophin stimulation for IVF, women with PCOS who are considering egg donation, immature oocytes collected for ART, use

of fresh or frozen ovarian tissues as sources of oocyte, IVM of germinal vesicle which is reconstructed¹.

In regards to different results of IVM in rates of oocyte maturation, fertilization, cleavage, and pregnancy^{38,39}, we designed this study to establish the relative success of unstimulated IVM in comparison with stimulated IVF in PCOS women.

Materials and Methods

During 2003, 70 PCOS women who were candidate for assisted conception were placed in two groups randomly: IVM and IVF. Finally, the prospective comparative study was done on 33 cycles of IVF. The patients commenced the treatment after full explanation and giving informed consent. An ovary was detected as PCO when showed ≥ 10 small (2-8 mm) cystic follicles around the periphery

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of ovary with enlarge echogenic stroma, during early follicular phase, using transvaginal ultrasound⁴⁰. Women with oligomenorrhea received IM injection of 100 mg of progesteron to induce withdrawal bleeding. In IVM group, a baseline ultrasound was done for all women on cycle days 3 and 8 to exclude dominant follicle and ovarian cysts, because data suggest that the presence of dominant follicle at the time of oocyte retrieval is deleterious to outcome of IVM¹². HCG priming, 10000 units, was done on days 9 to 12²⁶ and oocyte retrieval was done 36h after hCG injection with 18 g single-lumen aspiration needle with aspiration pressure of 7.5 kPa under general anesthesia. Oocytes were collected in culture tubes containing Hams F10 with 2 Iu/ml of heparin. After washing of blood, immature oocytes were incubated in a culture dish containing TC-199 medium supplemented with 25 mol/lit of pyruvic acid, 75 mIu/ml of FSH+LH, and 10% FBS at 37°C, 5% CO₂, and 95% air with high humidity. Maturity of oocytes was determined at 24 hours post collection denuded of cumulos. Mature oocytes were determined by presence of first polar body and ICSI was done and transferred into medium culture. Fertilization was assessed 18 hours after ICSI, with appearance of 2 pronuclei and 2 polarbodies. Embryo transfer was done on day 2 or 3 after ICSI. For endometrial preparation, Ethinyl estradiol was used depending on endometrial thickness on retrieval day and luteal support was provided by IM injection of 100 mg of progesteron/day.

In IVF group, pituitary suppression was done by long protocol with superfact (buserline) and

stimulation by hMG (225 unit/day), and serial control sonography was done to reveal mature follicles (at least 2 follicles with 18 mm in diameter). HCG priming with 10000 units of HCG was done and oocyte collection was done routinely and IVF or ICSI procedures were done traditionally. Outcome measure included the number of oocyte collected, metaphase II produced oocytes, embryos (2PN), pregnancy rate, and OHSS. In two groups statistical analysis were performed, using the SPSS statistical package. Chi square and fisher exact test were performed for statistical comparison between two groups.

Results

There were 35 women in IVM group and 35 in IVF. Two cycles in IVM group failed to reach egg retrieval. There were no differences in mean age, infertility cause, and previous IVF cycles. Oocytes retrieval in two groups were: 100% in IVF groups and 51.5% in IVM group ($P \sim 0$); fertilization rate of mature oocytes in IVM group was 30% per cycle and 100% per cycle in IVF group ($P \sim 0$); embryo transfer was 18.1% in IVM and 88.8% in IVF group ($P \sim 0$); pregnancy rate was 3.3% in IVM and 18.8% in IVF groups ($P \sim 0$); and OHSS was 11.2% in IVF groups and 0% in IVM group ($P < 0.05$) (Table 1).

Mean number of retrieved oocytes (mature or immature), fertilized oocytes, and transferred embryos in two groups showed significant differences (Table 2).

Table 1. Results of 33 unstimulated IVM and 35 stimulated IVF cycles in PCOS women.

	IVM	IVF	OR
Number of cycle	33	35	
Age	28.4	29.3	$P = 0.05$
Duration of infertility	8.2	7.5	$P = 0.51$
Oocyte collected (%)	17(51.5%)	35 (100%)	$P = 0.000$
Fertilized 2PN embryos (%)	10(30.3%)	35 (100%)	$P = 0.000$
Embryo transferred (%)	6(18.1%)	31(88.8%)	$P = 0.000$
Pregnancy rate (%)	1(3.03%)	4(11.4%)	$P = 0.357$
Moderation or sever OHSS (%)	0(0%)	4(11.4%)	

Table 2. Comparison of different outcome measures in two groups.

	IVM	IVF	P Value
Number of cycles	33	35	
Mean of retrieved oocytes per cycle	1.39	8.45	0.000
Mean of fertilized oocytes per cycle	0.27	3.85	0.000
Mean of transferred embryos per cycle	0.24	2.71	0.000

Table 3. Comparison of success rate in different stages cases with positive puncture in two groups.

Number	Positive puncture (percent)	Fertilized oocytes (percent)	Transferred embryos (percent)	Pregnancy rate (percent)
35	35 100%	35 100%	31 81%	4 (13%)
33	17 51%	10 60%	6 60%	1 16.6%
P value	0.000	0.000	0.03	61%

In table 3 we have compared the success rate in fertilization, and embryo transferring stages, and pregnancy rate in patients whom we succeed to retrieve oocytes.

Discussion

Laparoscopic electrocoagulation of the ovaries with PCOS can be an effective method for good response to stimulation, but adhesion formation results in 17% to 27% of patients. A similar outcome may be achieved by ovarian drilling by immature oocyte collection without adhesion. In addition, immature oocytes which retrieved by this method and used for IVM, can result in a good embryos.

Recent papers showed a high incidence (21%) of non cleavage embryos produced by IVM, and further (27%) failed to develop beyond the first division. In addition, the remaining embryos showed high rates of aneuploidy. Maybe, specific time and mediators affect from circulation and other follicles or cells around the immature oocyte came to expression or imprinting specific gene in vivo^{31, 34}. Therefore, it would be unwise to use these embryos in clinical setting until further

research. Early embryo development was controlled by maternal genome which inherited by oocyte genome. Therefore, gene expression and imprinting during oocyte maturation stages affect on embryo development⁴⁰. Both nuclear and cytoplasmic maturation need to ensure developmental competence of embryos. Synchrony between endometrial and embryonic development in fresh IVM ET isn't ideal⁴¹. The results of this study demonstrate that immature oocytes from PCOS patients could mature and fertilize invitro. Although number of oocytes which have been retrieved and number of embryos produced in this method were lower than other studies reported by Trounson et al⁹, Barnes et al²⁰, TJ Child², and Tan SL¹, but development in culture system and experience in oocyte retrieval from small follicles may provide an advantage for assisted reproductive in PCOS women in our center.

In regards to other studies^{19,1,2} in spite of our study, this protocol is a useful alternative treatment for PCOS women who are the candidates for ART.

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