

Effect of predictive value of progesterone level on the day of HCG injection for IVF success in women with infertility due to tubal factor or polycystic ovarian syndrome referred to the women hospital, Tehran, 2009

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Abstract

Background: Polycystic ovarian syndrome is one of the most common causes of endocrine disorders and main reason of infertility due to anovulation and recurrent abortions. Progesterone has been shown to have an important role in fertilization of oocyte and fetal implantation.

Objective: The purpose of this study was to compare the predictive value of progesterone level on IVF success in women with infertility due to tubal factor or PCOS.

Materials and Methods: In a stratified cohort study, we assigned 76 infertile women of 20-38 years old who referred to women hospital into two equal groups with fallopian tube factor infertility and PCOS. We measured the plasma levels of progesterone and estradiol on the day of HCG administration. The patients were divided into two groups based on progesterone level cut off point of 1.2ng/ml. Thereafter the incidence of pregnancy (chemical by β -HCG measurement and clinical by ultrasonography up to the 6 weeks after fetal transfer) was compared in these groups.

Results: Total pregnancy rates were 15.8% in patients with tubal factor infertility and 26.3% in women with PCOS. In women with PCOS, the pregnancy rate was less in patients with progesterone level <1.2 ng/ml. However this difference was not statistically significant. Likewise, we did not observe any significant differences in pregnancy rate in patients with fallopian tube factor infertility.

Conclusion: Serum progesterone level on the day of HCG administration is not well predictive of the IVF success in infertile women due to fallopian tube factor or PCOS. To obtain more uniform results, we recommend use of larger samples while the bias variable is taken into account and the ROC curve is used for determination of the unique serum progesterone level.

Key words: Serum progesterone Level, Infertility, PCOS, Tubal factor, IVF.

Introduction

An ovulatory dysfunction is a common problem and is responsible for about 40% of female infertility. Polycystic ovarian syndrome (PCOS) remains one of its leading causes (1). It is a main reason of infertility due to anovulation and recurrent abortions, which could be considered as a risk factor for breast cancer (1-2).

Clomiphene citrate is considered as the drug of choice for first line treatment of an ovulatory dysfunction (3), however, pregnancy

rate is disappointingly lower than expected (50% or less). Individuals not responding to this treatment are often called Clomiphene Citrate Resistant (CCR). An alternative medication for CCR's is traditionally administration of exogenous gonadotropine by injection. If this also fails the treatment for these cases is IVF (4).

Most often GnRH analogs are used in conjunction with gonadotropine to prevent the possibility of spontaneous LH surge and ovulation prior to the egg retrieval. It is reported that, despite an effective suppression

of endogenous gonadotropine by GnRH analog, there is a small increment in plasma progesterone up to 20% of stimulated cycles (5). Thus effect of progesterone increase on the cycle outcome is still controversial and not clear.

While some authors reported negative effect on the pregnancy rate when the plasma level of progesterone is more than 0.9 ng/ml on the day of HCG injection in IVF cycles (6), others such as Hoffman and coworkers (7, 8) did not find any significant relationship between these two items. They also stated that the high level of plasma progesterone on the day of HCG injection on ovum donors has no adverse effect on pregnancy rate. In 1993, Lergo *et al* (9) showed that premature LH surge with elevated progesterone level increased pregnancy rate of ovum donors during IVF cycles.

The aim of this study is evaluate the effect of plasma level of progesterone on the day of HCG injection on rate of pregnancy in IVF cycles of PCOS cases.

Materials and methods

This stratified cohort study was conducted in a period of one year (2009) on 76 infertile women with PCOS and tubal factor who were suitable candidates for the IVF program. The certification of medical ethic committee was approved by tehran university of medical sciences. All of these PCOS cases were between 20-38 years. They were evaluated for other causes of infertility with HSG, laparoscopy. Their partner's sperm analyses according to the guideline of WHO were all normal.

The criteria for selection of PCOS patients were based on: 1) A history of amenorrhoea or oligomenorrhoea, 2) Hyperandrogenism symptoms, eg. Hirsutism according to Ferriman Gallweg score that was greater than seven, 3) Elevated LH or LH/FSH >2, and 4) Increased ovarian volume more than 9 ml or antral follicle count of 10 or more in ultrasound imaging. Infertile women with tubal factor defined as obstruction of both tubes which documented by laparoscopy and have been candidate for IVF and aged between 20-38 years.

Excluding criteria were under 20 and over 38 years old, not candidates for IVF, other causes of infertility than tubal factor and PCOS, Male factor presence that needs TESE. All patients underwent long pituitary down regulation protocol. They were given low dose of OCP pill on the third day of menstruation then from 20-21 day of menstrual cycle (late luteal phase) GnRH agonist (Superfact Hocchest Aquila, Italy) 500 microgram per day sub cutaneous started.

After two weeks of administering superfact dose or the third day of next menstruation cycle, serum estradiol and vaginal sonographic scan were performed to confirm pituitary suppression. If serum estradiol level was below 50 Picogram/ml and there was no follicle or cyst greater than 10 mm and endometrium thickness was less than 4 mm induction ovulation would begin. Otherwise, GnRH agonist would be given for an extra week up to a maximum of 21 days.

150-225 unit gonadotropin [rFSH (gonal-f; Serono, Aubonne, Switzerland)] according to age, weight and FSH level on the third day of menstruation started. The superfact decreased to half of its initial dose on that day. After a period of 5 days of gonadotropin treatment, the dose of gonadotropin was regulated depending on ovarian response that was monitored by vaginal ultrasound. When there were at least two 18-mm follicles, HCG (Profasi, serono, Rome, Italy) 10000 IU was injected.

The progesterone level was checked on the day of HCG injection. Oocyte pickup was performed 36-38 h later. After a period of 48 hours, embryo transfer with Cook catheter (Cook catheter OB/GYN spencer JN) was carried out. The count of embryo transfer that depended on the age and quality of embryo was between 2-4. To support of luteal phase progesterone injection, 100 mg per day from the day of puncture of Oocyte was administered.

Experiment subjects were selected through interview, examination and lab tests. Upon receiving information, all subjects are consented to the experiment study. For each potential progesterone threshold from 0-1.8 ng/ml in increments of 0.1ng/ml. this method

is a variation of receiver operating characteristic (ROC) curve analysis (Honely and McNeil, 1982).

Serum progesterone breakpoints were optimized by summing the true positive rate and the true negative rate. According to serum progesterone on the day of HCG injection; the patients were divided to two groups based on progesterone level in HCG day. Clinical pregnancy defined as detecting of fetal heart by ultrasound at 6-8 week of gestation, and Chemical pregnancy rate was defined as only β -HCG<200 IU. The rate of pregnancy was compared between the two groups.

Statistical analysis

The experiment data were then compared against the Fisher's exact test in SPSS version 18. The level of significance was considered based on the criteria of $p < 0.05$.

Results

As stated earlier, 76 infertile women with PCOS and tubal factor who were suitable candidates for the IVF program. 38 patients were PCOS and 38 were tubal factor. Each group was divided to two clusters based on progesterone level on the day of HCG injection in IVF cycles. Overall pregnancy rate was 26.3% in PCOD patients and 15.8% in tubal factor group.

The findings of this experiment are summarized in Table I and II. Also 38 tubal factor patients joined in this study were divided to two groups based on progesterone level on the day of HCG injection in IVF cycles. The findings of this experiment are summarized in table III and IV.

Table I. Comparison of chemical pregnancy in PCOS patients according to progesterone level.

Group	Chemical pregnancy		Total
	No (%)	Yes (%)	
Progesterone <1.2	16 (84.21%)	3 (15.79%)	19 (100%)
Progesterone \geq 1.2	12 (63.16%)	7 (36.84%)	19 (100%)

Fischer`s Exact test p-value= 0.269.

Table II. Comparison of clinical pregnancy in PCOS patients according to progesterone level.

Group	Chemical pregnancy		Total
	No (%)	Yes (%)	
Progesterone <1.2	18 (94.74%)	1 (5.26%)	19 (100%)
Progesterone \geq 1.2	15 (78.95%)	4 (21.05%)	19 (100%)

Fischer`s Exact test p-value= 0.340.

Table III. Comparison of chemical pregnancy in tubal factor patients according to progesterone level.

Group	Chemical pregnancy		Total
	No (%)	Yes (%)	
Progesterone <1.2	16 (84.21%)	3 (15.79%)	19 (100%)
Progesterone \geq 1.2	16 (84.21%)	3 (15.79%)	19 (100%)

Fischer`s Exact test p-value= 0.99.

Table IV. Comparison of clinical pregnancy in tubal factor patients according to progesterone level.

Group	Clinical pregnancy		Total
	No	Yes	
Progesterone <1.2	17 (89.47%)	2 (10.53%)	19 (100%)
Progesterone \geq 1.2	17 (89.47%)	2 (10.53%)	19 (100%)

Fischer`s Exact test p-value=0.99.

Conclusion

The effect of progesterone level on the IVF outcome has been an uncertainty area for several years (10). Most studies used an absolute progesterone level on the day of HCG administration as an indicator of premature luteinization (PL) and the cutoff level differed from 0.8-2 ng/mL. Some authors defined PL as a progesterone/estradiol ratio of >1.

There is a marked variation in the incidence (13-71%), of PL due to discrepancies in definition, population characteristics and/or treatment protocols. The pathogenesis of PL in COH is not well understood. There are some hypotheses explain this subject: rise of follicular LH levels, serum accumulation of HCG from HMG, increased LH receptor sensitivity of the granulosa cells to FSH, or poor ovarian response with increased LH sensitivity. The role of this premature rise of serum progesterone on IVF outcome remain unclear (11).

In the present study, we have compared patients with PCOS and control patients (tubal factor) who had normal ovaries and assessed their pregnancy rate and effect of serum progesterone level on pregnancy rate. Total pregnancy rates were 26.3% in patients with fallopian factor infertility and 15.8% in women with PCOS. In women with PCOS, the pregnancy rate was less in patients with progesterone level <1.2 ng/ml. However this difference was not statistically significant. Likewise, we did not observe any significant differences in pregnancy rate in between two groups of patients with fallopian tube factor infertility.

Several authors did not find any negative effect of this on IVF outcome (12, 13). In two separate studies, Hoffman and coworkers (7, 8) observed no significant difference in pregnancy rate in patients undergoing IVF/ET with high or low progesterone concentration on the day of HCG administration and in patients who received oocytes donated from women with high or low progesterone concentrations. Huang *et al* (9) showed that there were no differences in clinical pregnancy

rate between 3 groups of patients with progesterone level less than 0.3 ng/ml, between 0.3 and 1 ng/ml, and more than 1 ng/ml on the day of HCG injection in IVF cycles.

Silverberg and coworkers (5) showed that increasing of serum progesterone level on the day of HCG injection had no adverse effect on the quality of Oocytes. Abuzeid *et al* (14) showed that level of progesterone on the day of HCG injection had no negative effect on pregnancy rate. Bustillo *et al* (15) cited that the level of progesterone on the day of HCG injection could not predict pregnancy rate. Lergo *et al* (8) presented PL with serum progesterone higher than 1.2 ng/ml increased pregnancy rate per embryo in donor cases. They also showed that PL with high level progesterone is a common event in IVF cycles.

It is a natural sign of follicular development; therefore it increases the pregnancy rate. Similarly, Doldi *et al* (4) showed premature increasing of serum progesterone in IVF PCOS cases could be a prediction factor of IVF success. On the contrary however, studies were found that related the PL to low pregnancy rate in IVF cycles. In their study, Schoolcraft and colleagues (16) showed serum progesterone greater than 0.5 ng/ml on the day of HCG injection had significant negative correlation with pregnancy rate in IVF cycles.

Harada *et al* (17) cited that small increase in serum progesterone on the day of HCG injection had inverse effect on IVF cycles. Burn *et al* (18) proved low pregnancy rate in IVF cycles with premature increase in serum progesterone. Drinfeild and collaborators (19) deduced that the pregnancy rate in IVF cycles was dropped if there was small increase in serum progesterone level on the day of HCG injection. Mio and coworkers (20) also found the same results. These findings suggest that PL may influence the endometrium, adversely affecting implantation and subsequent embryo development due to PL (21).

Based on this information and our results, it seems that premature progesterone production does not have an adverse effect on pregnancy rate, but on the contrary, may be a

predictor for success in IVF/ET. In conclusion, to obtain more uniform results, we recommend use of larger samples while the bias variable is taken into account and the ROC curve is used for determination of the unique serum progesterone level.

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References

1. Homburg R. Clomiphene citrate-end of an era? A mini review. *Hum Reprod* 2005; 20: 2043-2051.
2. Ghasemi N, Mortazavi MR, Khorasani A. Frequency of Poly cystic ovary syndrome in patients with premenopausal breast cancer. *Iran J Reprod Med* 2010; 8: 86-89.
3. Kamath MS, George K. Letrozole or clomiphene citrate as first line for anovulatory infertility. *Reprod Biol Endocrinol* 2011; 9: 86.
4. Doldi N, Marsiglio E, Destefani A, Gessi A, Merati G, Ferrari A. Elevated serum progesterone on the day of HCG administration in IVF is associated with a higher pregnancy rate in polycystic ovary syndrome. *Hum Reprod* 1999; 14: 601-605.
5. Edelstein MC, Seltman HJ, Cox BJ, Robinson SM, Shaw RA, Muasher SG. Progesterone levels on the day of human chorionic gonadotropin administration in cycles with gonadotropin-releasing hormone agonist suppression are not predictive of pregnancy outcome. *Fertil Steril* 1990; 54: 853-857.
6. Silverberg KM, Martin M, Olive DL, Burns WN, Schenken RS. Elevated serum progesterone levels on the day of human chorionic gonadotropin administration in in-vitro fertilization cycles do not adversely affect embryo quality. *Fertil Steril* 1994; 61: 508-513.
7. Hoffman GE, Bentzien F, Bergh PA, Garrisi GJ, Williams MC, Guzman I, et al. Premature luteinization in controlled ovarian hyper stimulation has no adverse effect on oocyte and embryo quality. *Fertil Steril* 1993; 60: 675-679.
8. Hoffman GE, Khoury J, Johnson CA, Thie J, Scott RT Jr. Premature luteinization during controlled ovarian hyper stimulation for in-vitro fertilization-embryo transfer has no impact on pregnancy outcome. *Fertil Steril* 1996; 66: 980-986.
9. Legro RS, Ary BA, Paulson RJ, Stanczyk FZ, Sauer MV. Premature luteinization as detected by elevated serum progesterone is associated with a higher pregnancy rate in donor oocyte in-vitro fertilization. *Hum Reprod* 1993; 8: 1506-1511.
10. Mio Y, Sekijima A, Iwabo T, Onohara Y, Harada T, Terakawa N. Subtle rise in serum progesterone during the follicular phase as a predictor of the outcome of in-vitro fertilization. *Fertil Steril* 1992; 58: 159-166.
11. Venetis CA, Kolibianakis EM, Papanikolaou E, Bontis J, Devroey P, Tarlatzis BC. Is progesterone elevation on the day of human chorionic gonadotropin administration associated with the probability of pregnancy in in vitro fertilization? A systematic review and meta-analysis. *Hum Reprod Update* 2007; 13: 343-355.
12. Elnashar AM. Progesterone rise on the day of HCG administration (premature Luteinization) in IVF: An overdue update. *J Assist Reprod Genet* 2010; 27: 149-155.
13. Martinez F, Coroleu B, Clua E, Tur R, Buxaderas R, Parera N, et al. serum progesterone concentration on the day of HCG administration cannot predict pregnancy in assisted reproduction cycles. *Reprod Biomed Online* 2004; 8: 183-190.
14. Huang JC, Jackson KV, Hornstein MD, Ginsburg ES. The effect of elevated serum progesterone during ovulation induction in in-vitro fertilization-embryo transfer. *J Assist Reprod Genet* 1996; 13: 617-624.
15. Abuzeid MI, Sasy MA. Elevated progesterone levels in the late follicular phase do not success of in-vitro fertilization- embryo transfer. *Fertil Steril* 1996; 65: 981-985.
16. Bustillo M, Stern JJ, Coulam CB. Serum progesterone at the time of human chorionic gonadotropin dose not predict pregnancy in in-vitro fertilization and embryo transfer. *Hum Reprod* 1995; 10: 2862-2867.
17. Schoolcraft W, Sinton E, Schlenker T, Huynh D, Hamilton F, Meldrum DR. Lower pregnancy rate with premature luteinization during pituitary suppression with leuprolide acetate. *Fertil Steril* 1991; 55: 563-566.
18. Harada T, Yoshida S, Katagiri C, Takao N, Ikenari T, Toda T, et al. Reduced implantation rate associated with a subtle rise in serum progesterone concentration during the follicular phase of cycles stimulated with a combination of a gonadotropin-releasing hormone agonist and gonadotropin. *Hum Reprod* 1995; 10: 1060-1064.
19. Burns WN, Witz CA, Klein NA, Silverberg KM, Schenken RS. Serum progesterone concentrations on the day after human chorionic gonadotropin administration and progesterone/ oocyte ratios predict in-vitro fertilization/ embryo transfer outcome. *J Assist Reprod Genet* 1994; 11: 17-23.
20. Dirnfeld M, Goldman S, Gonen Y, Koifman M, Lissak A, Abramovici H. A modest increase in serum progesterone levels on the day of human chorionic gonadotropin (hCG) administration may influence

pregnancy rate and pregnancy loss in in-vitro fertilization-embryo transfer (IVF-ET) patients. *J Assist Reprod Genet* 1993; 10: 126-129.

21. Edelman MC, Seltman HJ, Cox BJ, Robinson SM, Shaw RA, Muasher SJ. Progesterone level on the

day of human chorionic gonadotropin administration in cycles with gonadotropin-releasing hormone agonist suppression is not predictive of pregnancy outcome. *Fertil Steril* 1990; 54: 853-857.