



Comparison of Tamoxifen and Clomiphene Citrate for Ovulation Induction in Women with Polycystic Ovarian Syndrome: A Prospective Study

Sangita Sharma ^{1*}, Manisha Choudhary ¹, Vikas Swarankar ¹, Vaibhav Vaishnav ²

1- Department of Reproductive Medicine, Mahatma Gandhi University of Medical Science and Technology, Rajasthan, India

2- Manipal Hospital Jaipur, Rajasthan, India

Abstract

Background: The purpose of this study was to compare the efficacy of tamoxifen and clomiphene citrate in induction of ovulation in women with PCOS and anovulation.

Methods: In this prospective cohort study, 104 women with PCOS and primary infertility were enrolled after fulfilling the inclusion and exclusion criteria. The patients were allocated in two groups; group A (n=54) received tamoxifen 40 mg once daily (Days 3-7) and group B (n=50) received clomiphene citrate 100 mg once daily (Days 3-7). Serial ultrasounds were done till the administration of human chorionic gonadotropin (hCG). The ovulation and pregnancy rates in both groups were compared. The number of dominant follicles, estradiol levels, and endometrial thickness were also studied. Comparison was done using chi-square and student's t-test and a p-value of less than 0.05 was considered statistically significant.

Results: The number of dominant follicles and serum estradiol levels were significantly higher in group B (p<0.05), whereas the endometrial thickness was significantly more in group A (p<0.05). The ovulation rates were similar in both groups (66.6% vs. 70%, p=0.715). Pregnancy rate per treatment cycle and per ovulatory cycle was marginally higher in group A (14.81% and 22.22%, respectively), as compared to group B (14% and 20%, respectively), but the difference was not statistically significant (p>0.05).

Conclusion: Tamoxifen and clomiphene citrate are both equally effective in induction of ovulation and achieving a pregnancy in women with PCOS.

Keywords: Anovulation, Clomiphene, Infertility, Ovulation induction, Polycystic ovary syndrome, Pregnancy rates, Tamoxifen.

To cite this article: Sharma S, Choudhary M, Swarankar V, Vaishnav V. Comparison of Tamoxifen and Clomiphene Citrate for Ovulation Induction in Women with Polycystic Ovary Syndrome: A Prospective Study. *J Reprod Infertil.* 2021;22(4):274-281. <http://dx.doi.org/10.18502/jri.v22i4.7653>.

* Corresponding Author:
Sangita Sharma,
Department of Reproductive
Medicine, Mahatma Gandhi
University of Medical
Science and Technology,
Rajasthan, India
E-mail:
sangi237@yahoo.com

Received: Oct. 25, 2020
Accepted: Mar. 23, 2021

Introduction

About 10-15% of couples face the problem of infertility during their reproductive years (1). Ovulatory disorders are responsible for about 30 to 40% of female infertility (2). According to the WHO classification of anovulation, about 85% of anovulation cases fall under group II, which is hypothalamic pituitary dysfunction or eugonadotropic anovulation (3). This group consists predominantly of women with polycystic ovary syn-

drome (PCOS). Since the last six decades, ovulation inducing drugs have resulted in successful treatment of this group of infertile patients.

Clomiphene citrate (CC) has been widely used for ovulation induction in women with PCOS since its introduction in 1956 (4,5). The ovulation rate with clomiphene is around 80% (6), but still only 40% of these women are able to achieve a pregnancy (7). This discrepancy has been pro-

posed to be due to the antiestrogenic effects of clomiphene on cervical mucus (7) and endometrium (8). Tamoxifen is another non-steroidal selective estrogen receptor modulator (SERM) which has a similar mechanism of action like clomiphene, but acts as an agonist on the estrogen receptors of the endometrium and vaginal mucosa. It is commonly used as an adjuvant in the treatment of breast cancer. Its use in ovulation induction was first reported by Williamson and Ellis in 1973 (9). Although there is a wealth of data on clomiphene, the studies on the use of tamoxifen in ovulation induction are relatively inconclusive, with different studies reporting higher, lower or comparable results, when compared to clomiphene. However, some data is encouraging in terms of ovulation and pregnancy rates (9-11). Tamoxifen has given good results in women with clomiphene failure too (10). The reasons proposed for a better ovulation and pregnancy rates with tamoxifen are favorable cervical mucus and better endometrial thickness and functioning of corpus

luteum (10-14). The added advantages of tamoxifen are lower cost and lower chances of ovarian hyperstimulation. This study was conducted with the aim of comparing the efficacy of tamoxifen and clomiphene citrate in induction of ovulation in subfertile women with PCOS and anovulation.

Methods

This was a prospective cohort study, conducted over a period of one year in the Department of Reproductive Medicine of a tertiary care center in India. Sample size calculation was done with the use of the OpenEpi software. On the basis of the previous studies, to achieve a statistically valid comparison of the ovulation rates in both groups, considering type I error of 0.05 and a power of 80%, a sample size of at least 50 women in each group was required. One hundred and four women with polycystic ovary syndrome (Diagnosed by the Rotterdam's criteria) (15) and primary infertility were enrolled in the study after considering the inclusion and exclusion criteria (Figure 1). Proper

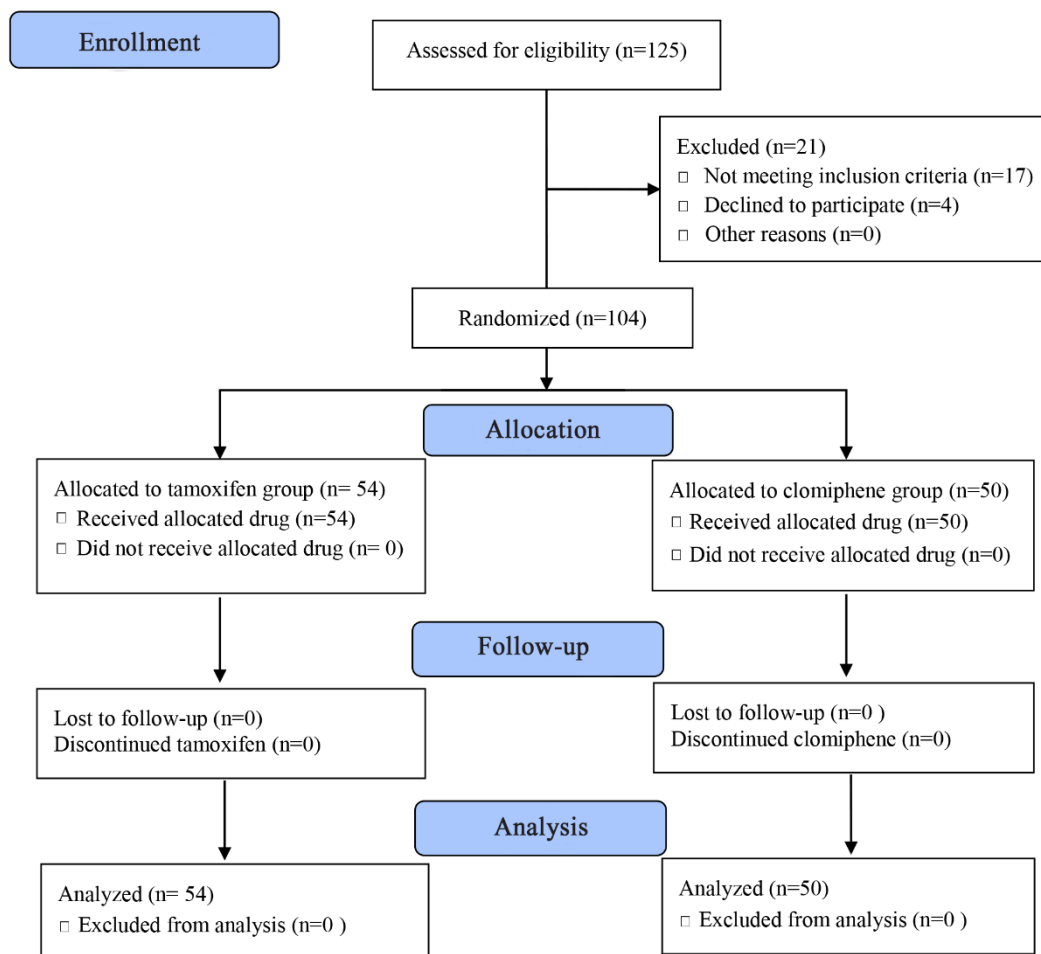


Figure 1. The study flow diagram

consent was taken from the patients and the study was approved by the institutional review board. Inclusion criteria of the study were (1) age of <35 years in anovulatory women, (2) BMI of 18-30 kg/m², (3) bilaterally patent fallopian tubes, and (4) semen parameters within normal range according to the 2010 WHO manual. Exclusion criteria of the study were (1) hydrosalpinx, adenomyosis or intramural fibroid >4 cm, (2) endometriomas seen in ovaries on ultrasound, (3) patients with previous history of ovarian drilling, (4) history of 3 previous unsuccessful ovulation induction cycles, and (5) other causes of anovulation (Primary or co-existing) such as hypothyroidism and hyperprolactinemia.

The hysterosalpingography showed bilateral spill of the radiopaque contrast from the fallopian tubes and the semen analysis report was also normal according to the 2010 WHO manual. On third day of a normal menstruation or withdrawal bleeding, serum FSH, LH and estradiol levels were measured, along with performing a baseline transvaginal scan. Although the included anovulatory patients were already diagnosed cases of PCOS, baseline serum FSH and LH levels were measured on the third day of the menstruation to compare the baseline characteristics in both groups. Once it was confirmed that the baseline endometrial thickness was less than 4 mm, there was no cyst in the ovaries and serum estradiol levels were less than 50 pg/ml, the patients were then allocated into group A (n=54) and group B (n=50) (Figure 1) by a nurse on the basis of a computer generated randomization table. The patients allocated in two groups; group A (n=54) received tamoxifen 40 mg once daily for 5 days (Days 3-7) and group B (n=50) received clomiphene citrate 100 mg once daily for 5 days (Days 3-7).

Serial ultrasounds were done from day 11 of the cycle to monitor the number and size of follicles, and to assess the endometrial thickness. Human chorionic gonadotropin (hCG) was administered intramuscularly at a dose of 5000 IU, when at

least one follicle reached a mean diameter of ≥18 mm on transvaginal scan. A repeat scan was done after 48 hr and ovulation was documented by one or more of the following criteria, viz. disappearance of the dominant follicle, decrease in the size of the follicle, change in its shape, appearance of internal echoes within it and/or appearance of free fluid in the pouch of Douglas. The couple was advised to have sexual contact during the periovulatory period after hCG administration. Luteal support was given by administration of micronized progesterone 200 mg twice daily vaginally for 14 days, after which serum beta hCG levels were measured. A transvaginal scan was done 2 weeks after the positive serum beta hCG report to evaluate the presence of gestational sac and confirm a clinical pregnancy. Pregnancies were followed till the 12th week of gestation.

Statistical analysis: The primary outcome measures were the number of developing dominant follicles in each patient (>16 mm), serum estradiol levels on the day of hCG trigger, endometrial thickness on the day of hCG trigger, and ovulation rate per treatment cycle. The secondary outcome measure was pregnancy rate per ovulatory cycle and also per treatment cycle. Comparison was done in both groups using chi-square and student's t-test and a p-value of less than 0.05 was considered statistically significant.

Results

Both groups were comparable in terms of age, BMI, basal FSH and LH, and duration of infertility (Table 1). The number of dominant follicles (>16 mm) was significantly more in the clomiphene group as compared to the tamoxifen group (2.1±0.2 vs. 1.02±0.4, p<0.05). On the day of hCG administration, the serum estradiol levels were significantly higher in the clomiphene group (196±21.4 vs. 168±15.6 pg/ml, p<0.05), whereas the endometrial thickness was significantly more in the tamoxifen group (10.4±0.3 vs. 8.8±0.7 mm, p<0.05). The ovulation rates were not significant-

Table 1. Patient characteristics in both the groups

	Tamoxifen (Gp A)	Clomiphene citrate (Gp B)	p-value
Number of women	54	50	
Age (Yrs)	26.4 (±2.8)	25.2 (±2.9)	0.921
BMI (kg/m ²)	26.2 (±3.6)	27.0 (±2.8)	0.734
Basal FSH (mIU/ml)	5.4 (±0.9)	4.9 (±1.2)	0.863
Basal LH (mIU/ml)	8.8 (±2.4)	9.2 (±1.01)	0.572
Duration of infertility (Yrs)	2.5 (±1.2)	2.7 (±1.6)	0.351

Values are expressed as Mean±SD

Table 2. Comparison of the results in tamoxifen and clomiphene groups

	Tamoxifen (Gp A) (n-54)	Clomiphene citrate (Gp B) (n-50)	p-value
No. of dominant follicles (≥ 16 mm)	1.02 (± 0.4)	2.1 (± 0.2)	0.001
Estradiol levels on the day of hCG administration (pg/ml)	168 (± 15.6)	196 (± 21.4)	0.023
Endometrial thickness on hCG day administration (mm)	10.4 (± 0.3)	8.8 (± 0.7)	0.006
Ovulation rate	36/54 (66.6%)	35/50 (70%)	0.715
Pregnancy rate per treatment cycle	8/54 (14.81%)	7/50 (14%)	0.905
Pregnancy rate per ovulatory cycle	8/36 (22.22%)	7/35 (20%)	0.818

Values are expressed as Mean \pm SD, absolute values (percentages in brackets)

ly different in both groups (Gp A-36/54 (66.6%), Gp B-35/50 (70%), $p=0.715$). Pregnancy rate per treatment cycle and per ovulatory cycle was marginally higher in the tamoxifen group (8/54 (14.81%) and 8/36 (22.22%), respectively), as compared to the clomiphene group (7/50 (14%) and 7/35 (20%), respectively), but the difference did not reach a statistical significance ($p=0.905$ and 0.818, respectively) (Table 2). There was one miscarriage in the tamoxifen group. There were no ectopic or multiple pregnancies in both groups.

Discussion

In the present study, the efficacy of tamoxifen for ovulation induction in subfertile women with PCOS was compared to that of clomiphene citrate. Not only the ovulation rates, but also the number of dominant follicles, peak serum estradiol levels, endometrial thickness, pregnancy rates per treatment cycle and per ovulation cycle were

compared and studied.

Ovulation rates: In our study, 36 women ovulated out of 54 (66.6%) in the tamoxifen group, whereas 35 women ovulated out of 50 (70%) in the clomiphene citrate group which showed no statistically significant difference between groups. These findings revealed higher ovulation rates as compared to the rates reported by Boostanfar et al. in a randomized control trial in which ovulation rates with tamoxifen and clomiphene citrate were 44.24 and 45.05%, respectively. The reason could be the difference in the daily dose of tamoxifen and clomiphene, which was 40 mg and 100 mg in our study and 20 mg and 50 mg in the RCT by Boostanfar et al., respectively. But interestingly, the ovulation rates with tamoxifen and clomiphene, in spite of being lower as compared to our study, were similar to each other in the RCT by Boostanfar et al. (17) (Table 3).

The ovulation rate with tamoxifen in our study

Table 3. Comparison of ovulation rates with tamoxifen and clomiphene citrate in different studies

Study	Year	Daily dose of tamoxifen (mg)	Daily dose of clomiphene (mg)	Ovulation rate with tamoxifen (%)	Ovulation rate with clomiphene (%)	p-value
Present study	2020	40	100	36/54 (66.6)	35/50 (70)	0.71 (NS)
Boostanfar et al. (RCT)	2001	20	50	50/113 (44.24)	41/91 (45.05)	>0.05 (NS)
Nardo (RCT)	2004	20	50	61/98 (62.24)	60/127 (47.24)	0.03 (In favor of Tamox)
Badawy and Gibreal (RCT)	2011	20	100	95/184 (51.63)	120/187 (64.17)	0.01 (In favor of CC)
Seyedoshohadaei et al. (RCT)	2011	10-30	50-150	34/50 (68)	39/50 (78)	>0.05 (NS)
Chunfeng and Musen (case-control study)	2016	20	50	31/38 (81.57)	25/38 (65.78)	<0.05 (In favor of Tamox)
Daqing et al. (RCT)	2016	20	50	42/49 (85.71)	33/49 (67.34)	<0.05 (In favor of Tamox)
Narayanan et al. (RCT)	2019	40-80	50-150	70%	71.4%	0.93 (NS)

was comparable to that reported by Nardo as 62.24% in a randomized control trial. Although the ovulation rate with clomiphene was much lesser (47.24%) as compared to the present study, the reason could again be the difference in the daily dose of clomiphene which was 100 mg in our study and 50 mg in the RCT by Nardo (18).

In 2011, an RCT conducted by Seyedshohadaei revealed ovulation rates of 68 and 78%, respectively in the tamoxifen and clomiphene groups, which was comparable to each other and similar to the present study (19).

In another RCT by Badawy and Gibreal in 2011, the ovulation rates were comparable to our study in the clomiphene group, 70 and 64.17%, respectively. In both studies, 100 mg of clomiphene was given daily. However, for tamoxifen, there was a difference in the dose, as it was 40 mg and 20 mg in our study and the RCT by Badawy and Gibreal, respectively. This explained the ovulation rates of 66.6 and 51.63%, respectively. Thus, Badawy and Gibreal reported significantly higher ovulation rates with clomiphene in comparison to tamoxifen, which was in contrast to the present study (20).

The ovulation rates with clomiphene in the present study were comparable to those reported by Chunfeng and Musen in a case-control study in 2016 and Daqing et al. in a RCT in the same year as 65.78 and 67.34%, respectively. However, the ovulation rates with tamoxifen were much higher in both of these studies, being 81.57 and 85.71%, respectively (21, 22). Moreover, Narayanan et al. reported comparable ovulation rates with tamoxifen and clomiphene (70% and 71.4%, $p=0.93$),

similar to the present study (23).

A meta-analysis of four trials, conducted by Steiner et al. in 2005, did not show any significant difference between the tamoxifen and the clomiphene groups regarding the chances of ovulation (OR 0.755, 95% CI 0.513–1.111) (24). Another recent meta-analysis, consisting of seven studies and 1573 participants, did not show any statistically significant difference in the ovulation rates in the tamoxifen and clomiphene groups (25).

Endometrial thickness: In the present study, the mean endometrial thickness was significantly higher in the tamoxifen group as compared to the clomiphene group (10.4+0.3 mm vs. 8.8+0.7 mm, $p<0.05$). This observation was similar to a few recent RCTs (20, 23, 26) and a case-control study (27) (Table 4). Badawy and Gibreal, Narayanan et al., and Sattar et al. found significantly higher mean of endometrial thickness with tamoxifen in comparison to clomiphene. However, the results of several studies are contradictory, showing similar endometrial thickness with both the drugs (25, 28-30).

Jie et al. in 2018 conducted a meta-analysis which included five studies (4 RCTs and one case-control study) to evaluate endometrial thickness on the day of hCG administration. No statistically significant difference was detected in the tamoxifen group when compared with the clomiphene group for endometrial thickness among anovulatory women (25). Interestingly, in most of the studies included in this meta-analysis, 50 mg clomiphene was used, whereas in those studies where endometrial thickness was significantly higher in the tamoxifen group as compared to

Table 4. Comparison of endometrial thickness with tamoxifen and clomiphene citrate in different studies

Study	Year	Daily dose of tamoxifen (mg)	Daily dose of clomiphene citrate (mg)	Mean endometrial thickness with tamoxifen±SD	Mean endometrial thickness with clomiphene citrate±SD	p-value
Present study	2020	40	100	10.4±0.3	8.8±0.7	<0.05
Badawy and Gibreal (RCT)	2011	20	100	10.1±0.1	9.3±0.4	<0.0001
Huang Zuekun et al. (case-control study)	2011	20	50	10.0±3.25	7.3±1.85	<0.05
Yu et al. (RCT)	2011	20	50	9.7±1.7	8.9±2.4	>0.05 (NS)
Lu (RCT)	2012	20	50	6.6±1.3	6.1±1.5	>0.05 (NS)
Lixia et al. (RCT)	2015	20	50	9.86±2.56	9.0±2.13	>0.05 (NS)
Narayanan et al. (RCT)	2019	40-80	50-150	10.4±0.45	8.77±0.96	<0.001
Sattar et al. (RCT)	2020	40	100	9.57±1.04	6.62±1.07	<0.001

clomiphene, the latter was given at the dose of 100 mg per day (20, 23, 26) (Table 4). It is known that higher doses of clomiphene results in better ovulation rates (6), but the antiestrogenic effect on endometrium is more with higher doses (31, 32); thus, the higher ovulation rates are not translated into higher pregnancy rates. With this background, along with comparable ovulation rates, and better endometrial thickness even at higher doses, tamoxifen seems to be a better alternative for ovulation induction in anovulatory women.

Pregnancy rates per cycle: The pregnancy rates per treatment cycle in our study were 14.81 and 14% for the tamoxifen and clomiphene group, respectively whereas the pregnancy rates per ovulatory cycle were 22.22 and 20% for tamoxifen and clomiphene groups. They were marginally higher in the tamoxifen group, but it was not statistically significant. Most of the studies have reported pregnancy rates per treatment cycle and very few have mentioned pregnancy rates per ovulatory cycle. The observations of the present study were

comparable to those of a few more studies in the recent past (17, 18, 28, 30) (Table 5).

Although the pregnancy rates in the clomiphene group in our study were similar to those reported by Chunfeng and Musen (21.05%), Daqing et al. (18.36%), and Qiu-yan (20%), the pregnancy rates reported by these authors in the tamoxifen group were much higher (42.10%, 38.77% and 45%, respectively) (21, 22, 33).

Similar to our study, the meta-analysis conducted by Steiner et al. in 2005 (Including four studies, with 273 women) also did not show any statistically significant difference in the pregnancy rates with tamoxifen and clomiphene (24). However, due to small sample size, the results were considered inconclusive and larger studies were required to be conducted.

A systematic review consisting of five trials comparing clomiphene with tamoxifen showed no clear evidence of a difference in live birth, clinical pregnancy, miscarriage, or multiple pregnancy rate (34).

Table 5. Comparison of pregnancy rates with tamoxifen and clomiphene citrate in different studies

Study	Year	Daily dose of tamoxifen (mg)	Daily dose of clomiphene citrate (mg)	Pregnancy rate per treatment cycle with tamoxifen (%)	Pregnancy rate per treatment cycle with clomiphene citrate (%)	p-value
Present study	2020	40	100	8/54 (14.8)	7/50 (14)	0.905 (NS)
Boostanfar et al. (RCT)	2001	20	50	10/113 (8.84)	6/91 (6.59)	>0.05 (NS)
Nardo (RCT) (pregnancy rate per ovulatory cycle)	2004	20	50	14/61 (22.9)	11/60 (18.3)	>0.05 (NS)
Badawy and Gibreal (RCT)	2011	20	100	20/184 (10.69)	35/187 (18.71)	0.04 (In favor of CC)
Seyedoshohdaei et al. (RCT)	2011	10	50	20/174 (11.49)	32/199 (16.08)	<0.05 (In favor of CC)
Yu et al. (RCT)	2011	20	50	8/40 (20)	8/44 (18.18)	>0.05 (NS)
Lixia et al. (RCT)	2016			17/80 (21.25)	16/80 (20)	>0.05 (NS)
Chunfeng and Musen (case-control study)	2016	20	50	16/38 (42.10)	8/38 (21.05)	<0.05 (In favor of Tamox)
Daqing et al. (RCT)	2016	20	50	19/49 (38.77)	9/49 (18.36)	<0.05 (In favor of Tamox)
Qiu-yan	2016	20	50	18/40 (45)	8/40 (20)	<0.05 (In favor of Tamox)
Narayanan et al. (RCT)	2019	40-80	50-150	15%	20%	0.32

A recent meta-analysis by Jie et al. including 10 studies and 1879 participants did not show any significant difference in the pregnancy rates when tamoxifen and clomiphene were compared (25).

In the present study, tamoxifen administration in comparison with clomiphene citrate resulted in few dominant follicles, better endometrial thickness, and comparable ovulation and pregnancy rates. The added advantages of tamoxifen can be lower incidence of ovarian hyperstimulation syndrome, cost effectiveness, and also a lower risk of epithelial ovarian cancer in comparison to clomiphene citrate (25, 35).

Conclusion

In the present study, in women with PCOS, ovulation induction with tamoxifen resulted in lesser dominant follicles, better endometrial thickness and similar ovulation and pregnancy rates when compared to clomiphene citrate. Thus we conclude that tamoxifen and clomiphene citrate are equally effective in induction of ovulation and achieving a pregnancy in women with PCOS.

Acknowledgement

We would like to offer gratitude to Dr. M.L. Swarankar for his constant inspiration. We take this opportunity to thank Dr. T.R. Sharma and Mrs K.K. Sharma, who helped in the final proof readings of the manuscript. We also thank all the women who participated in this study.

Funding: Nil.

Conflict of Interest

There is no conflict of interest to disclose.

References

1. Hamilton-Fairley D, Kiddy D, Watson H, Paterson C, Franks S. Association of moderate obesity with poor pregnancy outcome in women with polycystic ovary syndrome. *Br J Obstet Gynaecol.* 1999;99(2): 128-31.
2. Berek JS. *Novak's textbook of gynecology.* 13th ed. US: Williams & Wilkins; 2002. 1432 p.
3. NICE guidelines. *Fertility problems: assessment and treatment (CG156);* 2013. 51 p.
4. Greenblatt RB, Barfield WE, Jungck EC, Ray AW. Induction of ovulation with MRL/41, preliminary report. *JAMA.* 1961;178:101-4.
5. Wolf LJ. Ovulation induction. *Clin Obstet Gynecol.* 2000;43(4):902-15.
6. Gorlitsky GA, Kase NG, Speroff L. Ovulation and pregnancy rates with clomiphene citrate. *Obstet Gynecol.* 1978;51(3):265-9.
7. Gysler M, March CM, Mishell DR Jr, Bailey EJ. A decade's experience with an individualized clomiphene treatment regimen including its effect on the postcoital test. *Fertil Steril* 1982;37(2):161-7.
8. Eden JA, Place J, Carter GD, Jones J, Alaghband-Zadeh J, Pawson ME. The effect of clomiphene citrate on follicular phase increase in endometrial thickness and uterine volume. *Fertil Steril.* 1989; 73(2):187-90.
9. Williamson JG, Ellis JD. The induction of ovulation by tamoxifen. *J Obstet Gynecol Br Commonw.* 1973;80(9):844-7.
10. Borenstein R, Shoham Z, Yemini M, Barash A, Fienstein M, Rozenman D. Tamoxifen treatment in women with failure of clomiphene citrate therapy. *Aust N Z J Obstet Gynaecol.* 1989;29(2):173-5.
11. Gulekli B, Ozaksit G, Turhan NO, Senoz S, Oral H, Gokman O. Tamoxifen: an alternative approach in clomiphene resistant polycystic ovarian syndrome patients. *J Pak Med Assoc.* 1993;43(5):89-90.
12. Nicholas S, Macklon. *Optimizing protocols for ovulation induction. Female infertility therapy.* London: Martin Dunitz; 2000. p. 75-83.
13. Fox R, Corrigan E, Thomas PA, Hull MGR. The diagnosis of polycystic ovarian in women with menstrual disorder. *Fertil Steril.* 2000;66:761-4.
14. Roumen FJ, Doesburg HW, Rolland R. Treatment of infertile women with a deficient postcoital test with two antiestrogens: clomiphene and tamoxifen. *Fertil Steril.* 1984;41(2):237-43.
15. Rotterdam ESHRE/ASRM-Sponsored PCOS consensus workshop group. Revised 2003 consensus on diagnostic criteria and long-term health risks related to polycystic ovary syndrome (PCOS). *Hum Reprod.* 2004;19(1):41-7.
16. World Health Organization. *WHO laboratory manual for the examination and processing of human semen.* 5th ed. Geneva: World Health Organization; 2010. 271 p.
17. Boostanfar R, Jain JK, Mishell DR Jr, Paulson RJ. A prospective randomized trial comparing clomiphene citrate with tamoxifen citrate for ovulation induction. *Fertil Steril.* 2001;75(5):1024-6.
18. Nardo LG. Management of anovulatory infertility associated with polycystic ovary syndrome: tamoxifen citrate an effective alternative compound to clomiphene citrate. *Gynecol Endocrinol.* 2004; 19(50):235-8.
19. Seyedoshohadaei F, Zandvakily F, Shahgeibi S.

- Comparison of the effectiveness of clomiphene citrate, tamoxifen and letrozole in ovulation induction in infertility due to isolated unovulation. *Iran J Reprod Med.* 2012;10(6):531-6.
20. Badawy A, Gibreal A. Clomiphene citrate versus tamoxifen for ovulation induction in women with PCOS: a prospective randomized trial. *Eur J Obstet Gynecol Reprod Biol.* 2011;159(1):151-4.
 21. Chunfeng Sun, Musen Wang. Comparative study on the curative effect of clomifene and tamoxifen in infertile patients with polycystic ovary syndrome. *China Health Stand Manage.* 2016;13:121-3.
 22. Daqing Xiang, Ling Chen, Ling Zeng. Clinical efficacy of clomiphene and tamoxifen on infertility patients with polycystic ovary syndrome. *Iran J Reprod Med.* 2016;2:284-7.
 23. Narayanan M, Jahaan U, Gupta N. Comparative evaluation of different cost effective ovulation induction drugs and their effect on follicular growth, endometrial thickness and pregnancy outcome. *Int J Reprod Contracept Obstet Gynecol.* 2019;8(11):4549-53.
 24. Steiner AZ, Terplan M, Paulson RJ. Comparison of tamoxifen and clomiphene citrate for ovulation induction: a meta-analysis. *Hum Reprod.* 2005;20(6):1511-5.
 25. Jie L, Li D, Yang C, Haiying Z. Tamoxifen versus clomiphene citrate for ovulation induction in infertile women. *Eur J Obstet Gynecol Reprod Biol.* 2018;228:57-64.
 26. Sattar MM, El-Halaby AE, El-Shamy ES, Taha SN. Effect of clomiphene citrate, tamoxifen, and letrozole on endometrial thickness in cycles of ovulation induction: a randomized controlled trial. *Menoufia Med J.* 2020;33(2):405.
 27. Huang Xuekun Shang, Huiling Zhang, Siyou. Different stimulate ovulation drugs influence of early pregnancy outcome in patients with polycystic ovary syndrome. *Pract Med J.* 2011;27:3760-2.
 28. Yu Li, Dong-zi Yang. Letrozole, tamoxifen or clomiphene citrate for ovulation induction in women with polycystic ovarian syndrome after pretreatment: a prospective randomized trial. *Chin J Pract Gynecol Obstetr.* 2011;27:606-8.
 29. Lu Ye. Letrozole, tamoxifen and clomiphene citrate compared the clinical effect of treatment of polycystic ovary syndrome. *Chin Foreign Med Res.* 2012;10:11-2.
 30. Lixia Zhang, Li Yu. Three kinds of drugs in the treatment of polycystic ovary syndrome with infertility clinical comparative study. *Stud Chin Foreign Med.* 2015;32:57-9.
 31. Roumen FJ. [Decreased quality of cervix mucus under the influence of clomiphene: a meta-analysis]. *Ned Tijdschr Geneesk.* 1997;41(49):2401-5. Dutich.
 32. Gadalla MA, Huang S, Wang R, Norman RJ, Abdullah SA, El Saman AM, et al. Effect of clomiphene citrate on endometrial thickness, ovulation, pregnancy and live birth in anovulatory women: systematic review and meta-analysis. *Ultrasound Obstet Gynecol.* 2018;51(1):64-76.
 33. Qiu-yan Luo. The clinical effect of letrozole, tamoxifen and clomiphene in treatment of patients with polycystic ovary syndrome infertility. *Clin Med Res Pract.* 2016;1:39-40.
 34. Brown J, Farquhar C. Clomiphene and other anti-oestrogens for ovulation induction in polycystic ovarian syndrome. *Cochrane Database Syst Rev.* 2016;12(12):CD002249.
 35. Cook LS, Weiss NS, Schwartz SM, White E, McKnight B, Moore DE, et al. Population-based study of tamoxifen therapy and subsequent ovarian, endometrial and breast cancers. *J Natl Cancer Inst.* 1995;87(18):1359-64.