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Comparison of Effectiveness of Different Protocols used for Controlled Ovarian Hyperstimulation in Intra-Uterine Insemination Cycle

--Manuscript Draft--

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Abstract:	<p>Introduction: Intra-uterine insemination (IUI) is one of the most commonly performed procedure of assisted reproductive technologies (ART), for the treatment of infertility. Controlled ovarian hyperstimulation (COH) is an important first step while performing IUI. This study aims at establishing a relationship between stimulation protocol and pregnancy outcome following IUI.</p> <p>Methods: This is a retrospective study of 1001 cycles of IUI in which the patients were divided into two groups: Group A (CC only) and Group B (CC+ GN or GN alone). The primary outcome assessed was Clinical Pregnancy Rates (CPR) and the secondary outcomes were Miscarriage Rate (MR), Multiple Pregnancy Rates (MPR), Follicle Numbers and Endometrial Thickness (ET).</p> <p>Results: Significantly, higher CPR was observed in Group B in comparison to Group A (14.55% v/s 7.82%; p = 0.05). MR was much higher in Group A in comparison to Group B, (14.29% v/s 4.65%; p = 0.94), but it was non- significant. The Follicle Number and the Endometrial Thickness of the Group A v/s Group B are (1.54 ± 0.69 v/s 1.90 ± 1.04; p = 0.0003) and (8.56 ± 1.33 v/s 8.39 ± 1.29; p = 0.1784), respectively and for subgroups, Group B1 and Group B2 are 1.92 ± 0.99 v/s 1.65 ± 0.92; p = 0.0008 and 8.32 ± 1.27 v/s 8.69 ± 1.24; p = 0.0004), respectively.</p> <p>Conclusion: GN, either alone or the combination with CC, gives a higher CPR and a lower abortion rate following IUI, however, increasing the Multiple Pregnancy Rate.</p>

Title: Comparison of Effectiveness of Different Protocols used for Controlled Ovarian Hyperstimulation in Intra-Uterine Insemination Cycle

Short Title: Effects of Controlled Ovarian Hyperstimulation on Intrauterine Insemination Result

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Ethical statement:

The authors declare that there is no conflict of interest regarding the publication of this paper.

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12 reproductive technologies (ART), for the treatment of infertility. Controlled ovarian hyperstimulation (COH) is an
13
14 important first step while performing IUI. This study aims at establishing a relationship between stimulation protocol
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16 and pregnancy outcome following IUI.

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18 **Methods:** This is a retrospective study of 1001 cycles of IUI in which the patients were divided into two groups:
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20 Group A (CC only) and Group B (CC+ GN or GN alone). The primary outcome assessed was Clinical Pregnancy
21
22 Rates (CPR) and the secondary outcomes were Miscarriage Rate (MR), Multiple Pregnancy Rates (MPR), Follicle
23
24 Numbers and Endometrial Thickness (ET).

25
26 **Results:** Significantly, higher CPR was observed in Group B in comparison to Group A (14.55% v/s 7.82%; $p =$
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28 0.05). MR was much higher in Group A in comparison to Group B, (14.29% v/s 5.43%; $p = 0.94$), but it was non-
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30 significant. The Follicle Number and the Endometrial Thickness of the Group A v/s Group B are $(1.54 \pm 0.69$ v/s 1.90
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32 ± 1.04 ; $p = 0.0003$) and $(8.56 \pm 1.33$ v/s 8.39 ± 1.29 ; $p = 0.1784$), respectively and for subgroups, Group B1 and
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34 Group B2 are 1.92 ± 0.99 v/s 1.65 ± 0.92 ; $p = 0.0008$ and 8.32 ± 1.27 v/s 8.69 ± 1.24 ; $p = 0.0004$), respectively.

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36
37 **Conclusion:** GN, either alone or the combination with CC, gives a higher CPR and a lower abortion rate following
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39 IUI, however, increasing the Multiple Pregnancy Rate.

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41 **Keywords:** Infertility, Intrauterine Insemination, Clomiphene Citrate, Gonadotropins, Controlled Ovarian
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4 **Introduction**
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7 Infertility, a worldwide health issue is the inability of a couple to conceive, despite unprotected intercourse for the
8 period of one year. Most common causes for infertility include mild endometriosis, male factor, unexplained
9 infertility, ovulatory dysfunction, and aging (1, 2). Intrauterine insemination (IUI) is the most performed treatment in
10 women with infertility owing to mild male factor infertility, anovulation, endometriosis with at least one patent tube,
11 and unexplained infertility (3, 4, 5).
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16 Simplicity, easy management, low cost, and absence of potentially serious complications make IUI a leading
17 routinely used technique for infertility (3, 6, 7). Further, development in the type of stimulation protocols,
18 gonadotropins, ultrasound monitoring and sperm preparation techniques has enhanced the success rate of IUI.
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22 IUI may be performed in natural cycle as well as in combination with controlled ovarian hyperstimulation (COH)
23 (6). A higher CPR was observed in the latter cases in some studies (8), thus leading to using potential ovarian
24 stimulants such as clomiphene citrate (CC). Higher pregnancy rates are observed in patients who were given CC (5,
25 9). Bae and co-workers observed a higher rate of pregnancy in patients administered with 3-days treatment of CC
26 than those given 5-day treatment with CC. This was believed to be due to an adequate endometrial growth by CC in
27 the 3-Day treatment (9). Gonadotropins (GN) such as human menopausal gonadotropin (hMG), follicle stimulating
28 hormone (FSH) have also been used as an alternative source for COH (6, 10, 11, 12). GNs although more responsive
29 in COH than CC, are expensive and have certain associated AEs [Ovarian Hyperstimulation Syndrome (OHSS), and
30 high-order multiple pregnancies]. But, being highly responsive and effective these are used in COH (5). Using a
31 combination of CC and GN; COH and IUI can be made a cost effective and efficient process.
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43 In this study, we assessed the efficacy and safety of CC, GN alone and in combination with CC for COH in infertile
44 female patients undergoing IUI. The study aimed at establishing a standard ovulation stimulation protocol including
45 the type and dose of the ovarian stimulants (including CC and GNs *etc.*) to be used combined with IUI to treat
46 infertility.
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51 **Materials and Methods**
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53 This was a retrospective, observational study conducted from 1 November, 2014 to 29 February, 2016 under good
54 clinical practices as per the International Conference on Harmonization (ICMR, 2006) guidelines and the Declaration
55 of Helsinki (13). Women unable to conceive due to ovulatory dysfunction, mild male factors, unexplained infertility,
56 mild endometriosis, *etc.* were included in the study. Following the physical examination and medical history
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4 collection, baseline scan for the female patients (willing to participate) was performed on Day 2 or Day 3. The
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6 patients were divided into two groups: Group A received CC and Group B received GN [Human Menopausal
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8 Gonadotropin (hMG)]. The patients in Group B were subdivided into two subgroups: Group B1 received GNs (hMG)
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10 plus CC and Group B2 received GN only. **Figure 1** represents the patient disposition. The dosage details are given in
11
12 **Figure 2.**

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14 Post semen collection by masturbation and liquefaction (at room temperature), it was analysed for pre wash count
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16 and motility under microscope. The concentrate of motile sperms was obtained using either swim up or density
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18 gradient technique and was re-analysed for post wash count. A final volume of 0.5 ml was used for insemination.

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20 IUI was performed 38 hours after the administration of Injection hCG (Injection Ovitrelle® 250 microgram, Merck
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22 Sereno) using the treated semen samples. Urine pregnancy test was performed to confirm the positive pregnancy if
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24 the patients did not have their periods within two weeks and if positive, sonography was performed one week later.
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26 The CPR was the primary outcome, while miscarriage rate (MR), multiple pregnancy rate (MPR), endometrial
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28 thickness and follicle number were the secondary outcomes. These were compared among different patient groups
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30 using SPSS version 19.0 (IBM Corporation, United States).
31

32 33 **Results**

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35 A total of 1001 patients were distributed into three different groups depending on the type of the treatment given.
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37 **Table 1** gives the number of patients with different etiologies for infertility among the two patient groups; which was
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39 found to be non-significant ($p = 0.1161$).

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41 CPR varied significantly between Group A ($n=9$) and Group B ($n=129$). It was higher in Group A v/s Group B
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43 (7.82% v/s 14.55 %); ($p = 0.05$). The subgroup analysis of Group B1 ($n=101$) and Group B2 ($n=28$) showed
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45 comparable CPR *i.e.* (14.44% v/s 14.97%, respectively); ($p = 0.85$) (**Table 2**).

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47 MR was higher in Group A patients ($n=1$) in comparison to Group B ($n=7$) (14.29% v/s 5.43%; $p = 0.94$); but the
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49 difference was not significant. The test of significance could not be performed in the groups B1 (6/129, 4.65%) and
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51 B2 (1/129, 0.077%) due to a small sample size.

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53 While no multiple pregnancies were observed in Group A, the MPR was 13.17 % in Group B. Among a total 17
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55 patients exhibiting multiple pregnancy; 11 and 6 patients belonged to Group B1 and B2, respectively, which was
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57 non-significant (10.89% v/s 21.42%; $p = 0.14$). Thus, MPR was higher in patients receiving GN alone.
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4 Group A had a lower number of follicles than Group B (1.54 ± 0.69 v/s 1.90 ± 1.04 ; $p = 0.0003$) which was found to
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6 be statistically significant, while Group B1 had a higher number of follicles than Group B2 (1.92 ± 0.99 v/s $1.65 \pm$
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8 0.92 ; $p = 0.0008$). The difference in endometrial thickness was non-significant between groups (Group A: $8.56 \pm$
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10 1.33 v/s Group B: 8.39 ± 1.29 ; $p = 0.17$). The comparison between the subgroups, showed that the follicle number
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12 (1.92 ± 0.99 v/s 1.65 ± 0.92 ; $p = 0.0008$) was significantly higher in Group B1 than in Group B2, respectively.
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14 While, endometrial thickness (8.32 ± 1.27 v/s 8.69 ± 1.24 ; $p = 0.0004$) was significantly higher in Group B2
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16 **(Table3).**

17 18 **Discussion**

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20 The effectiveness of different protocols for COH that can be used in combination with IUI was assessed in the
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22 current study in terms of primary (CPR) and secondary outcomes (MR, MPR, Follicle Numbers, Endometrial
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24 Thickness).

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26 In the present study, CPR was significantly higher in Group B patients who received GN alone or in combination
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28 with CC, in comparison to Group A patients, *i.e.*, 14.55 % and 7.82% ($p = 0.05$) who received CC alone. CPR was
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30 comparable between the groups (Group B1 v/s Group B2) receiving GN+CC or GN alone (14.44% v/s 14.97%; $p =$
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32 0.86).

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34 Previous studies have reported varying outcomes on following use of CC and hMG alone and in combination.
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36 Deshpande *et al* (2013) reported 13.89%, 15.58% and 7.43% CPR, in patient groups treated with hMG + CC, hMG
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38 only and CC only, respectively (7). The pregnancy rate was higher when low dose of hMG (75 IU) was administered
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40 either alone or in combination with CC (15.58% and 13.89%, respectively). Similar conclusion was drawn by
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42 Kamath *et al* (2010) (10).

43
44 The per cycle pregnancy rates for CC+IUI, FSH+IUI and IVF were 7.6%, 9.8% and 30.7%, respectively, obtained
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46 during FASTT (The Fasttrack And Standard Treatment Trial) study (14). Recently, Karadag *et al* (2016) observed a
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48 higher CPR in the GN group than in the CC group (15). The higher pregnancy rates after using GN could be due to a
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50 better follicular development, development of multiple follicles and good endometrial thickness.

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52 Recently, in a study conducted by Diamond *et al* (2015) in women with unexplained infertility aged 18-40 years, the
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54 CPRs in the patient groups given GN alone v/s CC alone were 35.5% v/s 28.3%, respectively. The multiple gestation
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56 rates were higher in GN only group (31.8%; $p = 0.006$) than that in CC only group (9%; $p = 0.44$). Mostly, twins (4,
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58 1.3%) were found to be born in CC only group, while there were 25 twins (8.3%) and 6 (2%) triplets in GN only
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4 treatment group (16). This could possibly be due to multiple follicular development and multiple ovulation. The
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6 present study also have demonstrated multiple follicles with the use of GNs.

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8 Mukherjee and coworkers conducted a prospective, randomized study comparing the effect of the single dose of
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10 uFSH + CC with only CC. The CPR observed was 17% v/s 8.3% ($p = 0.0001$) in patient groups administered with a
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12 combination of uFSH + CC v/s CC only, respectively. The difference in MR was observed ($p = 0.99$). MR was
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14 higher in Group A patients ($n=1$; 14.29%) than Group B ($n= 7/129$; 5.43%); but the difference was not significant (p
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16 = 0.94). The test of significance could not be performed in the groups B1 (6/129; 4.65%) and B2 (1/129; 0.077%)
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18 due to a small sample size. Higher MR in CC group can be due to inappropriate Endometrial Thickness due to its
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20 anti-oestrogenic nature.

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22 In an open labelled, randomized study by Peerear *et al* (2015), CPR in hMG only v/s CC only patient groups were
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24 14.4% (48/334)v/s 9% (29/323).The same study reported 8.5 mm v/s 7.5 mm endometrial thickness in hMGv/s CC
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26 patients groups, respectively ($p < 0.001$).The CPR and Endometrial Thickness in our study were 14.97% v/s 7.82%
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28 and 8.69 ± 1.24 mm v/s 8.56 ± 1.33 mm in the GN (hMG) only (Group B2)v/s CC (Group A) only groups,
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30 respectively. Thus, the results were in agreement with the study by Peerear and coworkers.

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32 Denkart and coworkers compared the CPR, MPR and live births in patient groups (with unexplained infertility and
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34 male subfertility) given CConly and FSH only. The respective outcome rates were 38% and 34.3%, 7.4% and 4.3%
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36 and 70.4% and 73.9%, in CC only and FSH only groups, respectively(17). The dose given in the current study was
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38 almost same, for all the treatment groups, as in the above two studies, *viz.*, GN alone group was given 75 IU daily
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40 from Day 2/3 of natural cycle; CC+GN group was administered with 50 mg CC from Day 2- Day 6 and 75 IU GN on
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42 alternate days as Day 3, Day 5 and Day 7. The comparable rates for the outcomes assessed were observed in the
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44 above two studies and the present study.

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46 Karadag *et al* observed no multiple pregnancies in both the patient groups (15). However, in the present study, no
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48 cases of multiple pregnancies were observed in CC alone patient group, while a higher rate was seen in hMG alone
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50 group (21.42%) than in CC+hMG group (10.89%).

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52 However, in a study by Abdelazim and co-workers showed contrasting results to the present study wherein CPR was
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54 higher in CC+hMG v/s hMG only group; 26.7% v/s 6.7%, respectively. Although, no significant differences were
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56 found in the outcomes such as endometrial thickness and number of follicles (18). Whereas, in the present study
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58 significant differences were observed in the endometrial thickness (8.35 ± 1.44 mm v/s 8.63 ± 1.43 mm in
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4 CC+hMGv/shMG only group, respectively) and number of follicles (2.01 ± 0.99 v/s 1.69 ± 0.94 in CC+hMGv/shMG
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6 only group, respectively) for the concerned groups.

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8 Thus, based on the findings of present study and the past research, we could state that the protocol followed in our
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10 study could be used as an effective, standard protocol for COH with IUI. The stepwise protocol given in **Figure 2**
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12 should be followed.
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14 **Conclusion**

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16 GN, either alone or in combination with CC, gives a higher CPR and a lower MR following IUI, however, increasing
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18 the MPR. Thus, GN or the combination of GN and CC can be used for an effective COH in combination with IUI for
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20 improved CPRs.
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References

- (1) Kumar S, Mohsen N, Malini S. Association of Obesity with Male Infertility among Infertile Couples is not Significant in Mysore, South India. *Adv Stud Biol* 2013;5: 319-325.
- (2) Shadap A. Causes of Infertility among Married Women-A Review.
- (3) Soria M, Pradillo G, García J, Ramón P, Castillo A, Jordana C, Paricio P. Pregnancy predictors after intrauterine insemination: analysis of 3012 cycles in 1201 couples. *J Reprod Infertil* 2012;13: 158-166.
- (4) Merviel P, Heraud MH, Grenier N, Lourdel E, Sanguinet P, Copin H. Predictive factors for pregnancy after intrauterine insemination (IUI): an analysis of 1038 cycles and a review of the literature. *Fertil Steril* 2010;93: 79-88.
- (5) Chaudhary K, Suri V, Dhaliwal LK, Gainer S. Comparison of the efficacy of letrozole and low-dose gonadotropin combination with clomiphene and low-dose gonadotropin combination as a controlled ovarian stimulation regime prior to intrauterine insemination in patients with unexplained infertility. *Fertility Science and Research* 2014;1: 98.
- (6) Koli P, Anil M, Ramya N, Patil K, Swamy M. Intrauterine insemination: a retrospective review on determinants of success. *Int J Reprod Contracept Obstet Gynecol* 2013;2: 311-314.
- (7) Deshpande SA, Deshpande AB. Intra uterine insemination an experience in rural population. *IJMRHS* 2013;2: 341-349.
- (8) Liu J, Chian R, Ma X, Wang W, Cui Y. The Pregnancy Outcomes Comparison on Natural or Controlled Ovarian Stimulation Cycles in Intrauterine Insemination Treatment: An Analysis of 8,893 Cycles. *La Prensa Medica* 2016;2016:
- (9) Bae S-A, Joo J-K, Choi J-R, Kim S-S, Lee K-S. Clinical outcomes of three-or five-day treatment with clomiphene citrate combined with gonadotropins and a timed intercourse cycle in polycystic ovary syndrome patients. *CERM* 2015;42: 106-110.
- (10) Kamath MS, Bhave P, Aleyamma T, Nair R, Chandy A, Mangalaraj AM, Muthukumar K, George K. Predictive factors for pregnancy after intrauterine insemination: A prospective study of factors affecting outcome. *J Hum Reprod Sci* 2010;3: 129.

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3
4 (11) Kaser DJ, Goldman MB, Fung JL, Alper MM, Reindollar RH. When is clomiphene or gonadotropin
5
6 intrauterine insemination futile? Results of the Fast Track and Standard Treatment Trial and the Forty and
7
8 Over Treatment Trial, two prospective randomized controlled trials. *Fertil Steril* 2014;102: 1331-1337. e1331.
9
10 (12) Rashidi BH, Tanha FD, Rahmanpour H, Ghazizadeh M. Luteal phase support in the intrauterine insemination
11
12 (IUI) cycles: a randomized double blind, Placebo Controlled Study. *J Family Reprod Health* 2014;8: 149.
13
14 (13) Hurst SA. Declaration of Helsinki and protection for vulnerable research participants. *JAMA* 2014;311: 1252.
15
16 (14) Reindollar RH, Regan MM, Neumann PJ, Levine B-S, Thornton KL, Alper MM, Goldman MB. A
17
18 randomized clinical trial to evaluate optimal treatment for unexplained infertility: the fast track and standard
19
20 treatment (FASTT) trial. *Fertil Steril* 2010;94: 888-899.
21
22 (15) Karadag B, Dilbaz B, Karcaaltincaba D, Sahin EG, Ercan F, Karasu Y, Tonyali NV. The effect of luteal-phase
23
24 support with vaginal progesterone on pregnancy rates in gonadotropin and clomiphene citrate/intra-uterine
25
26 insemination cycles in unexplained infertility: A prospective randomised study. *J Obstet Gynaecol* 2016;36:
27
28 794-799.
29
30 (16) Diamond MP, Legro RS, Coutifaris C, Alvero R, Robinson RD, Casson P, Christman GM, Ager J, Huang H,
31
32 Hansen KR. Letrozole, Gonadotropin, or Clomiphene for Unexplained Infertility. *N Engl J Med* 2015;373:
33
34 1230-1240.
35
36 (17) Dankert T, Kremer J, Cohlen B, Hamilton C, Pasker-de Jong P, Straatman H, van Dop P. A randomized
37
38 clinical trial of clomiphene citrate versus low dose recombinant FSH for ovarian hyperstimulation in
39
40 intrauterine insemination cycles for unexplained and male subfertility. *Hum Reprod* 2007;22: 792-797.
41
42 (18) Abdelazim IA, Makhlof HH. Sequential clomiphene citrate/hMG versus hMG for ovulation induction in
43
44 clomiphene citrate-resistant women. *Arch Gynecol Obstet* 2013;287: 591-597.
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Table and Figure Legends

Table 1: Etiology of Infertility for Group A (CC) and Group B (CC+GN or GN only)

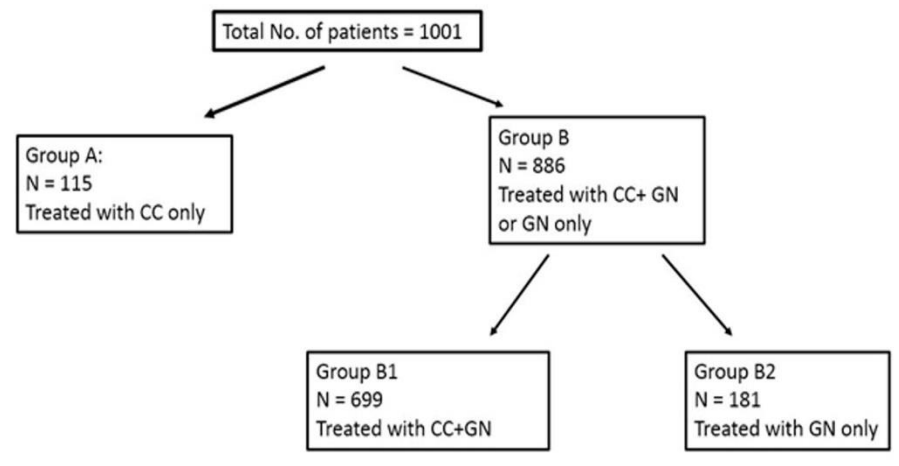
Table 2: Primary and Secondary Outcomes Obtained for Group A (CC) and Group B (CC+GN /GN only)

Table 3: Primary and Secondary Outcomes Obtained for Group B1 (CC+GN) and Group B2 (GN only)

Figure 1: Patient Groups as per the Treatments Given in the Present Study

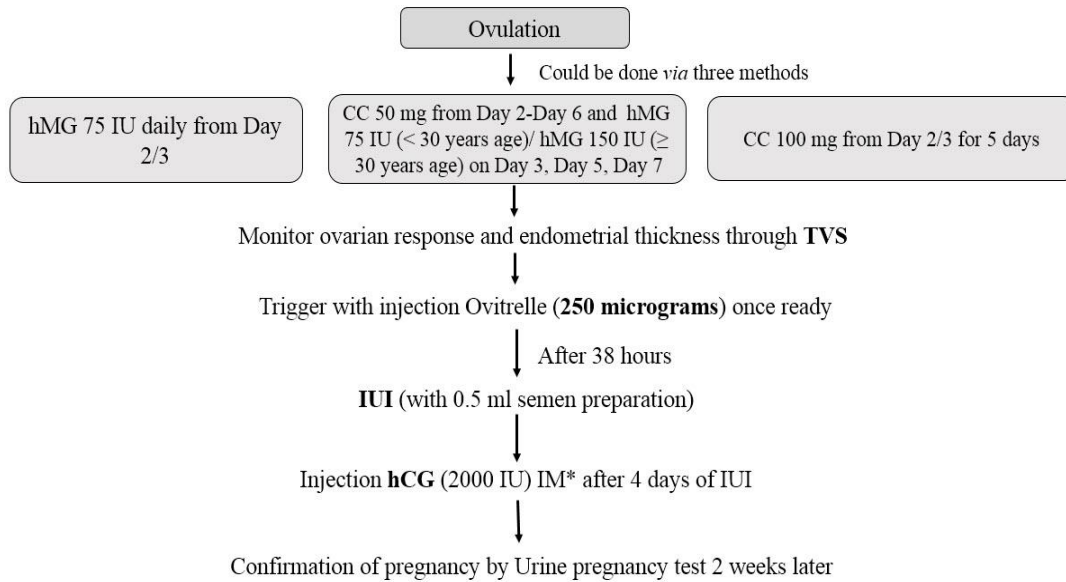
Figure 2: Illustration of the Ovarian Stimulation Protocol Standardized During the Present Study

Figure 1: Patient Groups as per the Treatments Given in the Present Study



CC: Clomiphene citrate; GN: Gonadotropins; N: total number of patients in a particular group

Figure 2: Illustration of the Ovarian Stimulation Protocol Standardized During the Present Study



*Administer hCG (10000 IU) when the endometrial thickness reaches ≥ 7 mm and follicle diameter reaches ≥ 18 mm.

CC- Clomiphene citrate; hMG: Human Menopausal Gonadotropin; FSH: Follicle Stimulation Hormone; IU: International Units; TVS: Transvaginal Ultrasonography; hCG: Human Chorionic Gonadotropin; IUI: Intrauterine Insemination; IM: Intramuscular.

Table 1: Etiology of Infertility for Group A (CC) and Group B (CC+GN or GN only)

Etiologies	CC (n = 115)	CC+GN or GN only (n = 886)
Endometriosis (n = 56)	6 (0.05%)	50 (0.06%)
Male factor (n = 233)	36 (0.31%)	197 (0.22%)
Multifactorial (n = 68)	11 (0.09%)	57 (0.06%)
Ovulatory dysfunction (n = 242)	22 (0.19%)	220 (0.25%)
Unexplained infertility (n = 402)	40 (0.35%)	362 (0.41%)

p-value: 0.1161

Table 2: Primary and Secondary Outcomes Obtained for Group A (CC) and Group B (CC+GN /GN only)

Outcome Rates (%)	Group A (N= 115)	Group B (N= 886)	<i>p</i> value
Primary Outcome	9/115 (7.82%)	129/886 (14.55%)	0.05
Clinical Pregnancy			
Secondary Outcome (Mean ± SD)			
Multiple Pregnancy	0	17/129 (13.17%) [^]	
Miscarriage Rate	1/9 (14.29 %)	7/129 (5.43%)	0.94
Follicle number	1.54 ± 0.69	1.90 ± 1.04	0.0003
Endometrial thickness (mm)	8.56 ± 1.33	8.39 ± 1.29	0.1784

^{*}Group 2A and Group 2B are subgroups of Group B[^] non-evaluable *p*-value.

Table 3: Primary and Secondary Outcomes Obtained for Group B1 (CC+GN) and Group B2 (GN only)

Outcome Rates (%)	Group B1 (N= 699)	Group B2 (N= 187)	<i>p</i> value
Primary Outcome Clinical Pregnancy	101/699 (14.44%)	28/187 (14.97%)	0.85
Secondary Outcome (Mean ± SD)			
Multiple Pregnancy	11/101 (10.89%)	6/28 (21.42%)	0.14
Miscarriage	6/129 (0.047%)	1/129 (0.008%)	
Follicle number	1.92 ± 0.99	1.65 ± 0.92	0.0008
Endometrial thickness (mm)	8.32 ± 1.27	8.69 ± 1.24	0.0004

CC+GN: 699 and GN only: 187

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