Mid Trimester Termination of Pregnancy: Role of Combined Mifepristone and Misoprostol

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Abstract

Background: Mid trimester abortions constitute 10%-15% of all induced abortions worldwide. Over the last decade this increase is due to better prenatal screening. It can be done by surgical and medical methods. Medical methods such as Misoprostol is widely used for mid trimester abortion. Mifepristone has antiprogesterone properties, so addition of Mifepristone with Misoprostol can increase its effectiveness. To assess the safety, effectiveness and acceptability of combined Mifepristone and Misoprostol for mid trimester medical termination of pregnancy (Between 13-24 weeks of gestation).

Materials and methods: This experimental study was conducted among 40 healthy women who presented for mid trimester termination of pregnancy between 13-24 weeks with missed abortion, gross congenital anomalies with or without previous history of one caesarian section. The study was conducted from March October 2018 at Chattogram Maa-O-Shishu Hospital Medical College, Chattogram, Bangladesh. Each woman received a single dose of tablet Mifepristone 200mg. After 24 hours, 200 mcg vaginal Misoprostol was administered which was repeated at 6 hourly interval for maximum of 5 doses (1000 mcg) in 24 hours. Success was taken as complete expulsion of fetus and placenta within 24 hours of first dose of Misoprostol. Primary and secondary outcomes were measured. Statistical analysis was done using SPSS version 23.

Results: Success rate of complete abortion was 97.5%. Mean Induction Abortion Interval was 11.59 hours (SD± 3.34). Mean dose of Misoprostol was 1.85 (SD± 0.77) or 370 mcg. Overall safety of the study was satisfactory with only 1 patient experienced fever and 1 had nausea. There was no major complication.

Conclusion: The Mifepristone/Misoprostol regimen is a highly effective as well as safe option for mid trimester medical termination of pregnancy with a short induction abortion interval and it can also be used in scarred uterus with close supervision.

Key words: Mifepristone; Misoprostol; Mid trimester abortion; Medical termination of pregnancy.

INTRODUCTION

Mid trimester abortion constitutes 10%-15% of all abortion but are responsible for two-third of all major complications (WHO 1997)1. There has been a gradual increase in the second trimester termination of pregnancy because of the widespread introduction of prenatal screening programs which detects serious fetal anomalies2. Various surgical and medical methods have been tried for the second trimester MTP with varying success and induction abortion interval. During the last decade medical
methods for second trimester induced abortions have been considerably improved. According to the WHO and Royal College of Obstetrics and Gynaecologists Mifepristone followed by Misoprostol is considered as an effective and safe method for second trimester abortions\(^3,4\). Prostaglandin (PGE1, PGE2 and PGF2alpha) has been used for mid trimester pregnancy termination in the last 20 years. When prostaglandin E1 analogue (Gameprost or misoprostol) are used alone for second trimester MTP, the mean Induction Abortion Interval (IAI) can be as long as 12-16 hours\(^2,5\). Mifepristone (RU486 a substitute 19-norethisterone derivative) acts by blocking progesterone receptors. Progesterone is a key hormone in maintaining pregnancy by keeping the uterus in a quiescent state. It prevents softening and dilatation of the cervix, reduces prostaglandin output from the deciduas and suppresses uterine contractions. So, pre treatment with antiprogesterone prostaglandins required as well as the analgesia requirement\(^5,7\). The combination of Mifepristone and Misoprostol has been found safe and effective for mid trimester termination of pregnancy in various studies\(^4,8\).

The present study was conducted to assess the effectiveness of combination of Mifepristone followed by successive doses of vaginal misoprostol for mid trimester abortions and to study any side effects of the above regimen.

**METHODS AND MATERIALS**

This prospective study was done in Chattagram Maa-O-Shishu Hospital Medical College from March October 2018. A total of 48 patients presented to us for termination of pregnancy between 13-24 weeks period of gestation. 40 eligible patients were included in the study after taking written informed consent and proper counseling in accordance with the inclusion criteria Ethical approval was taken from the ethical committee of the institute.

**Inclusion criteria**

i) All patients admitted in the labour ward seeking for medical termination for missed abortion or major congenital anomalies

ii) Patients with history of 1 caesarian section were also included in the study.

**Exclusion criteria**

i) Women who were hemodynamically unstable at the time of presentation

ii) Women who had either taken MTP pill from outside or self-prescribed, came with inevitable or incomplete abortion

iii) Women with known case of heart disease, uncontrolled hypertension, bronchial asthma or coagulation disorder, on anti coagulant or corticosteroid therapy

iv) Haemoglobin <9gm%

v) > 1 caesarian section.

All women were given tab Mifepristone (200 mg) to be swallowed under supervision. After 24 hours, tab Misoprostol (200 mcg) was administered per vaginum in the posterior fornix which was repeated at 6 hourly intervals for maximum 5 doses (1000 mcg) for 24 hours depending on cervical dilatation, uterine contraction and history of previous scar. Sterile vulval pads were given which were examined at regular intervals. Onset of uterine contractions, vaginal bleeding and passage of any product of conception and placental completeness were monitored. The time of expulsion of fetus and placenta was noted. When placenta was not expelled spontaneously, evacuation and curettage was done. Successful termination was considered as the occurrence of complete abortion within 24 hours of first dose of Misoprostol.

**Following Primary Outcomes were Measured:**

i) Rate of complete abortion after initial Misoprostol administration

ii) Induction abortion interval (IAI) calculated from the time of first dose of Misoprostol administration to complete expulsion of fetus and placenta

iii) Mean dose of Misoprostol required.

**Secondary Outcomes were:**

i) Side effects like nausea/ vomiting, diarrhoea, fever, headache

ii) Any complications observed with the regimen like excessive bleeding, incomplete abortion, need for blood transfusion, sepsis, scar rupture or uterine injury.

**RESULTS**

A total of 48 women underwent mid trimester pregnancy termination during the study period. Out of these, 4 patients came with inevitable abortion, 2 had self administered abortion pills and 2 were with previous history of 2 caesarian section so they were excluded from the study. Finally 40 cases were included in the study and analysed.

The mean age of the women included in the study was 23 years with a range of 18-32 years. 32.5%of total patients were nullipara, 22.5% were Primigravida and remaining 45% were multipara. 17.5%women presented in 13-16 weeks,55% were in 17-20 weeks and 27.5% of women were in 21-24 weeks of gestation. 32.50% women came with previous history of abortion. 3 women came with previous caesarian section and 3 women came for MTP for congenital malformations eg. neural tube defect.

**Table1:** Demographic profile of mothers (n=40)

| Particular                        | Frequency (n) | Percentage (%) |
|----------------------------------|--------------|----------------|
| Age (Years)                      |              |                |
| 1. <19 years                     | 03           | 7.50%          |
| 2. 19-24 years                   | 22           | 55.00%         |
| 3. 25-29 years                   | 11           | 27.50%         |
| 4. >29 years                     | 04           | 10.00%         |
| Parity                           |              |                |
| 1. 0                             | 13           | 32.50%         |
| 2. 1                             | 09           | 22.50%         |
| 3. >=2                           | 18           | 45.00%         |
Nullipara (n=13) Primipara (n=9) Multipara (n=18)

Successful complete abortion (Expulsion of fetus and placenta) 24 hours after the instillation of first dose was found in 39 cases (97.5%). However, evacuation was done in 1 case due to incomplete expulsion. There were 3 cases of previous history of 1 caesarian section. All of them were presented in 17-20 weeks of gestation. In these group of patients mean induction abortion interval was 10.83 hours. All of these patients had complete abortion and none required surgical intervention.

9 patients received injectable analgesics for moderate discomfort. 7 out of these were primigravida. Oral analgesics were sufficient for pain relief in 21 patients. There were 10 patients who did not require any analgesics.

Table 4: Side effects (n=40)

There was no major complications observed. Only 1 patient had fever and 1 patient had nausea. None had diarrhea or uterine rupture. There was no case of major obstetric haemorrhage observed during the study.

DISCUSSION

Mid trimester MTP is a difficult situation owing to the prolonged time required for the abortion process and associated complications. In developing countries especially rural areas second trimester MTP is a real challenge owing to the limited resources available. In earlier days surgical methods used be the standard method for mid trimester MTP. In the last 20 years there is emergence of medical methods for mid trimester pregnancy termination consisting of prostaglandins alone or in combination with anti-progesterone Mifepristone. Still a method which is 100% reliable, safe and affordable is not known.

Misoprostol as an effective abortificient is being successfully used through all routes i.e sublingual, oral, vaginal and in different regimens with varying induction abortion interval. Combination of Mifepristone and Misoprostol with different dose schedules is widely used method for 2nd trimester MTP. Priming of the uterus with Mifepristone makes it more sensitive to prostaglandin. It binds with the progesterone receptors and antagonizes the action of progesterone resulting in increase in prostaglandin production and decreased deactivation of prostaglandins. It also softens the cervix, thus enhancing the efficacy of prostaglandin.

We have given Misoprostol 200 mcg vaginally 6 hourly for 5 doses 24 hours after priming with 200 mg of oral Mifepristone. It has been seen that vaginal route of administration of Misoprostol is safer and more effective than oral route with less side effects due to better bioavailability of the drug at the target site9,10.
We compared our study with other studies where Mifepristone 200mg followed by minor variations of subsequent dosages and routes of administration of Misoprostol. Various studies have shown higher success rate and reduced induction abortion interval and need for lesser dose of Misoprostol when Mifepristone is added to Misoprostol\(^5,7,11\). Gupta et al\(^{10}\) gave the second dose of Misoprostol vaginally or sublingually after 4-6 hours of first dose\(^{12}\). The success rate was 91.42% compared to that of ours where the success rate is 97.5%. Ashok et al used vaginal route for the first dose and oral for the subsequent doses of Misoprostol and the success rate was 97.1% which is very close to our success rate\(^5\).

In our study, minimum IAI is 5 hours 15 minutes, maximum IAI is 19 hours and mean induction abortion interval is 11.59 hours which is very close to the another study performed by Maninder et al following the same regimen and the induction abortion interval was 11.26 hours\(^{13}\). Some other studies like Ashok PW et al and Goh et al the induction abortion interval were 6.5 hours and 6.7 hours respectively though they followed different dose schedules and various routes of Misoprostol administration after 200 mg of Mifepristone intake\(^5,14\).

In our study mean number of dose of Misoprostol was found to be 1.85 and the total dose was calculated to be 370 mcg which is also close to the study by Maninder et al the mean total number of dose was 2.13 and mean total dose was 610.42 mcg\(^{13}\). Our present study is also comparable to other studies\(^5,13\).

None of the patients in our study aborted completely with Mifepristone alone though studies have reported 0.2%-0.5% incidence of complete abortion with only Mifepristone\(^5,8,12\).

There were 3 patients in our study who had undergone caesarean section prior to present pregnancy. All of them had complete abortion with combination of Mifepristone and Misoprostol. Ther were various case reports showing uterine rupture in previously scarred uterus undergoing mid-trimester pregnancy termination\(^{15}\). Again many have shown safety of Mifepristone and Misoprostol for mid trimester MTP in cases of previous caesarian section\(^{16,17}\).

In our study, there was no relation found between the parity and induction abortion interval which is consistent with the study performed by Gupta et al\(^{12}\).

There was no case of rupture uterus or need for hysterotomy. Nulliparous patients required more injectable analgesics as compared to the multiparous women which can be due to their low threshold for pain which also shows the similarity with the literature\(^2,9\).

LIMITATION

The limitations of the present study were its sample size and not using the randomization.

CONCLUSION

Use of mifepristone and misoprostol is highly effective and safe option for mid-trimester medical termination of pregnancy. It has shorter induction abortion interval, high success rate and less side effects with significantly short hospital stay. It can also be used in scarred uterus with caution. However, a broad based, well designed randomized study is hereby suggested for national policy formulation.

DISCLOSURE

All the authors declared no competing interest.
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