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Original Research Article

An observational study to evaluate fall in hemoglobin following abortions conducted by misoprostol and surgical methods

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ABSTRACT

Background: Over the past three decades, medical methods of abortion have been developed throughout the world and are now the standard methods of providing abortion care in addition to surgical methods. There are various differences between the two methods. In this study, evaluating the fall in haemoglobin levels in abortions conducted by misoprostol and D and E in women with pregnancy failure.

Methods: Total of 80 women who fulfilled the criteria were included in the study. Inclusion and exclusion criteria applied and Hb levels noted on day 1 and 15 during follow up. In addition to this induction abortion interval and side effects of both the methods were also studied. Participants assigned to medical treatment received 800 μg of misoprostol inserted into posterior vaginal fornix i.e., day 1st and repeat dose on day 3 if no expulsion. Follow up done after 1 week and 15 days, if there was no expulsion, suction and evacuation was done. Participants assigned to dilatation and evacuation group would undergo the procedure in operation theatre. Statistical analysis done.

Results: The mean induction to abortion interval in the misoprostol group was 9.1 ± 2.1 hours (mean \pm SD). The mean hemoglobin level in the misoprostol group on day 15 was 9.7 ± 1.12 and that in the D and E group was 10.26 ± 1.31 p value (0.04), statistically significant.

Conclusions: Proper counseling of the side effects in both the methods and prompt action to alarming signs are required to avoid major disaster.

Keywords: Abortion, D and E, Misoprostol

INTRODUCTION

In India, the second most populous country in the world, abortion has been legal on a broad range of grounds since 1971. Over the past three decades, medical methods of abortion have been developed throughout the world and are now the standard methods of providing abortion care. Medical abortion, which involves the use of medications rather than a surgical procedure to induce an abortion, is an option for women who wish to terminate a first trimester pregnancy. The method is most commonly used up to 63 days of gestation but is also effective after. The centres for disease control and prevention estimates that 64% of abortions are performed before 63 days of gestation. Medical methods are safe, effective and avoids invasive procedures as in surgical abortion but requires

active patient participation and follow up. Surgical methods on the other hand are completed in a single setting, have higher success rate but carries risk of perforation, adhesion formation and may affect the reproductive potential of women. In this study we are evaluating the fall in haemoglobin level in both the methods in women with early pregnancy failure.

Objective of this study was to evaluate the fall in haemoglobin following abortions conducted by misoprostol and D and E.

METHODS

This is an observational study carried out at Government Medical College Ambikapur from October 2019 to

February 2020. Total 80 women diagnosed with early pregnancy failure in ultrasound findings were included in the study.

USG findings included

- A crown-rump length ≥6 mm and no cardiac activity or
- A crown-rump length <6 mm and no fetal growth at least 1-week later or
- A gestational sac with absent embryonic pole for at least 1 week.

A written and informed consent was taken and inclusion and exclusion criteria were applied.

Inclusion criteria

- Gestational age of 12 weeks or less.
- Haemoglobin >9 g/dl
- Not allergic to prostaglandins.

Exclusion criteria

- Gestational age >12 weeks,
- Haemoglobin < 9 g/dl,
- Scarred uterus
- Currently on anti-coagulants
- Allergic to prostaglandins.
- Excessive vaginal bleeding.

Women were randomly assigned in the medical method or surgical methods of abortion flipping a coin with 40 participants in each group. Age, period of gestation, parity noted along with Hb concentration.

Participant's assigned to medical treatment received 800 μ g of misoprostol inserted into the posterior vaginal fornix on the day of enrolment, i.e., day 1st. If expulsion was not complete by day 3rd, they received a second dose of misoprostol (800 μ g) and follow up done after one week during which if there was no expulsion, suction and evacuation was done. Participants assigned to dilatation and evacuation group would undergo the procedure in operation theatre.

Both the groups were asked to come back on day 15 during which USG was done to look for retained products of conception in those who underwent medical methods of abortion and haemoglobin concentration checked. The criterion for successful medical treatment was the diagnosis of complete abortion, established by an endometrial thickness ≤15 mm. Medical treatment was considered to have failed if, after the first or second dose of treatment, it was not possible to achieve complete uterine evacuation and a new curettage was needed to complete the process. Associated factors like pain, bleeding, nausea, dizziness, fever was compared in both the groups.

Statistical analysis

Data were analyzed and statistical analysis done. Chi square test was used for analyzing the differences between the two proportions (p <0.05 was considered significant).

RESULTS

A total of 150 women who underwent ultrasound were diagnosed to have early pregnancy failure out of which 120 fulfilled the criteria but only 80 agreed to participate in the study and hence included.

Most of the women in both the groups belonged to 23 to 27 years of age, lower socioeconomic status, and gravida 2 with gestational age of 8 to 10 weeks upon admission. There was no significant difference among the two groups regarding age, parity or period of gestation. The (mean \pm SD) age of participants in the medical method was 25 \pm 4.9 years and that in the surgical method was 25 \pm 3.2 years (p value =1.0000).

About 37.5% were gravida 2 in both the groups whereas primigravida were 30% in the misoprostol group and 25% in the surgical group. Multigravida comprised 12.5% in each group.

Analyzing gestational age of the cases it was found that 12 (30%) and 15 (37.5%) were in 6-8 weeks gestation in misoprostol and D and E group, respectively; 20 (50%) were in 8-10 weeks gestation in the both groups respectively and the rest 8 (20%) in misoprostol and 5 (12.5%) in surgical group with 10-12 weeks of gestation. The clinical parameters are described in Table 1.

Table 1: Clinical characteristics of participants.

Parameters	Medica	al method	Surgical	method		
Age (in years)	N	%	N	%		
18-22	10	25%	7	17.5%		
23-27	16	40%	18	45%		
28-32	8	20%	12	30%		
>32	6	15%	3	7.5%		
Obstetric history						
G_1	12	30%	10	25%		
G_2	15	37.5%	15	37.5%		
G_3	8	20%	10	25%		
G ₄ or more	5	12.5%	5	12.5%		
Gestational age (in weeks)						
6-8	12	30%	15	37.5%		
8-10	20	50%	20	50%		
10-12	8	20%	5	12.5%		

The mean induction to abortion interval in the misoprostol group was 9.1±2.1 hours (mean±SD). Eighteen patients (45%) had aborted within 6 to 8 hours times span whereas 14 (35%) had expulsion within 9 to 10 hours while the remaining 8 (20%) had expulsion at

more than 11 hours. For those who had induction abortion interval more than 11 hours had gestational age >10 weeks (Figure 1).

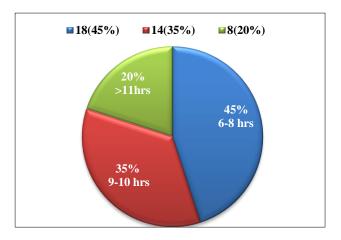


Figure 1: Induction abortion interval.

It was also noted that there was statistically significant difference between the general complaints of the women in both the groups. There was nausea in 20 (50%) participants in the misoprostol group whereas only 6 (15%) had complaints of nausea in the D and E group.

Dizziness was present in 10 (25%) and 8 (20%) in the misoprostol and D and E group respectively. Eighteen women (45%) in the misoprostol group had vomiting in contrast only 2 (5%) had vomiting in the D and E group. A total of 10 (25%) had complaints of diarrhoea and 8 (20%) had fever in the misoprostol group, on the contrary none such complaints were noted in the surgical methods. Pain and bleeding were also more common among the misoprostol group (Table 2).

Estimation of haemoglobin was done on day 1 as well as day 15 in both the groups. One participant in the misoprostol group had severe bleeding and required immediate transfusion hence excluded from the study. Another two women were lost to follow up while one required evacuation. Therefore, the total number of participants in the misoprostol group was 36 on day 15 of the study. The mean haemoglobin level in the misoprostol group on day 15 was 9.7±1.12 and that in the D and E group was 10.26 ± 1.31 p value (0.04), statistically significant. There was much fall in haemoglobin levels in the misoprostol group owing to minor episodes of bleeding throughout the follow up time duration and delay in seeking medical care. Women in the D and E group only six had bleeding till 15 days; rest of them had no such complaints (Table 3).

Complaints	Misoprostol n (%)	D and E n (%)	Statistical results
Nausea	20 (50%)	6 (15%)	Chi square = 11.028, DF=1, p value 0.0009 (significant)
Dizziness	10 (25%)	8 (20%)	Chi square = 0.283, DF=1, p value 0.59 (not significant)
Vomiting	18 (45%)	2 (5%)	Chi square = 16.853, DF=1, p value 0.0001 (significant)
Diarrhoea	10 (25%)	0 (0%)	Chi square = 4.158, DF=1, p value 0.041 (significant)
Fever	8 (20%)	0 (0%)	Chi square = 8.778, DF=1, p value 0.003 (significant)
Pain	30 (75%)	2 (5%)	Chi square = 40.323, DF=1, p value <0.0001 (significant)

Table 2: General complaints of participants.

Table 3: Haemoglobin levels (day 1 and day 15).

6 (15%)

Hb levels Misoprostol group		roup	D and E group		
(g/dl)	Day 1 (n=40) n (%)	Day 15 (n=36) n (%)	Day 1 (n=40) n (%)	Day 15 (n=40) n (%)	Statistical results
8-9.9	4 (10%)	10 (27.7%)	4 (10%)	4 (10%)	Chi square = 3.904, DF=1, p=0.04 (significant)
10-10.9	20 (50%)	22 (61.1%)	22 (55%)	26 (65%)	Chi square = 0.12, DF=1, p=0.72
11-11.9	12 (30%)	4 (11.1%)	10 (25%)	8 (20%)	Chi square = 1.114, DF=1, p=0.29
12-12.9	4 (10%)	0 (0%)	4 (10%)	2 (5%)	Chi square = 2.02, DF=1, p=0.15

DISCUSSION

Bleeding

28 (70%)

Several studies suggest that management of early pregnancy failure with misoprostol provides an effective alternative to D and E. Published reports emphasize success rates of medical management but provide little information about associated side effects including bleeding and fall in Hb levels. Expected bleeding patterns are an important part of patient counseling and may influence acceptability and choice of medical versus surgical management.

Chi square = 24.4, DF=1, p value <0.0001 (significant)

The most common pattern in the misoprostol group was some bleeding or spotting every day for 15 days. The median number of bleeding or spotting days after treatment was 12. In this analysis, heavy bleeding included the following categories: moderate, heavy or more than two pads/hours. Heavy bleeding was less frequent than light bleeding or spotting. Participants reported a median of 3 days of heavy bleeding. This is one of the major drawbacks in the medical methods leading to the decline in Hb levels.

In this study, the fall in mean hemoglobin level in the misoprostol group on day 15 was 9.7±1.12 and that in the D and E group was 10.26±1.31 p value (0.04), statistically significant. In the study conducted by Prema K, Ramalakshmi BA, while comparing Hb status in mifepristone versus vaccum aspiration in 303 women.² Hb estimations done after (40 days) abortion showed that there was no alteration in the mean Hb levels either in first or second trimester patients. This is contrary to this finding.

In other study conducted by Thonneau P et al, the women who had used the drug combination experienced a mean fall of 0.7 g/dl in haemoglobin (36% lost >1 g/dl and 8% >2 g/dl); haemoglobin concentrations remained stable in women who had vacuum aspiration.³ This is similar to this study.

Another study done by Davis et al with 800 μg vaginal misoprostol (repeat dose on day 3^{rd} of 800 μg if no evacuation) in missed abortion $\leq \! \! 13$ weeks gestation showed decreases in haemoglobin levels were greater after misoprostol (-0.7 g/dl) than curettage (-0.2 g/dl) (p $<\! 0.001$).⁴

In the present study 36 (90%) cases in misoprostol group underwent successful evacuation without resorting to surgical intervention. The mean induction to abortion interval was 9.1 ± 2.1 hours (mean \pm SD). In a study conducted by Thomas B et al in JIPMER in patients with missed abortion less than 14 weeks with 400 µg misoprostol used vaginally showed a mean insertion expulsion time 9.7 ± 5.6 hours. Out of 60 patients (76.7%) expelled the products completely. The present study also showed a similar result. In a similar study done by Singh K et al. In women with less than 8 weeks gestation with 800 µg vaginal misoprostol followed by a repeat dose of 400 µg every 3 hours for a maximum of 3 doses, found a complete evacuation rate of 84.7% with a mean time interval of 8.1 hours from initial.

The side effects associated with misoprostol are similar to that in other studies.^{7,8} Only one case in this study required transfusion due to heavy bleeding. Rests of the side effects were minor.

CONCLUSION

Misoprostol is an effective drug for induction of abortion where lesser of expertise is required in comparison to surgical methods. The main complaints is hemorrhage associated, which when not taken care may lead to adverse effects on the health of the women. Hence appropriate patient counselling as well as prompt action is the backbone which may decrease the associated morbidity.

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