Prophylactic Use of Intraumbilical Vein Oxytocin in the Management of Third Stage of Labor

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Abstract

**Background:** The third stage of labor is defined as the period of time between delivery of the fetus and delivery of the placenta. The most common complication accompanying this stage is postpartum hemorrhage (PPH) and prolonged third stage of labor owing to placenta retention and uterine atony are among the underlying cause of most cases of PPH. The duration of the third stage of labor is 5-15 min.

**Aim:** To assess the efficacy of intraumbilical vein oxytocin in reducing duration of third stage of labor and the amount of blood loss.

**Patients and Methods:** This study included 150 women attending the delivery room of Department of Obstetrics and Gynecology which was divided according to the inclusion and exclusion criteria into two groups:

- **Group A** (study group): include 75 cases which received 10 IU (1ml) of oxytocin in umbilical vein.
- **Group B** (control group): include 75 cases which received 1ml of saline in umbilical vein.

**Results:** The time of third stage took seconds in each group with no statistically significant difference detected (p>0.05).

**Conclusion:** Intra-umbilical oxytocin is a useful alternative in patients where methylergometrine is contraindicated or in cases where intravenous fluids need to be restricted. For optimum effect, rapid injection immediately after clamping of the cord is essential.

**Keywords:** intraumblical oxytocin, third stage of labour, postpartum hemorrhage

Introduction

The third stage of labor is defined as the period of time between delivery of the fetus and delivery of the placenta. The most common complication accompanying this stage is postpartum hemorrhage (PPH) and prolonged third stage of labor owing to placenta retention and uterine atony are among the underlying cause of most cases of PPH. The duration of the third stage of labor is 5-15 min (1).

Postpartum hemorrhage is a major cause of maternal death worldwide. It is the cause of approximately half of all maternal deaths in developing countries (2), and even in many developed countries. It is one of the major causes of admission of mothers to intensive care units (3).

The common causes of postpartum hemorrhage include bleeding from the site of implantation of the placenta, genital and nearby organs trauma or both of them (4).

Hemostasis of placental site is first established with the contraction of myometrium and thrombosis of the vessel lumens. As a result, parts attached to the placenta or large blood clots which impede efficient contracting of myometrium, can disrupt hemostasis in the placental site (5).

Aim of the Work

This is a randomized control study aims to assess the efficacy of intraumbilical vein oxytocin in reducing duration of third stage of labor and the amount of blood loss.

**Patients and Methods**

**Type of Study:** Randomized controlled study.

**Study Settings:** Al-Galaa Teaching Maternity Hospital.

**Study Duration:** 6 months (from April 2017 – September 2017).

**Study Population:**

This study included 150 women attending the delivery room of Department of Obstetrics and Gynecology which was divided according to the inclusion and exclusion criteria into two groups:

- **Group A** (study group: include 75 cases which received 10 IU (1ml) of oxytocin in umbilical vein).
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- Group B (control group: include 75 cases which received 1ml of saline in umbilical vein).

The inclusion criteria:
1. Uncomplicated a live single pregnancy.
2. 37 to 41 weeks gestational age.
3. Vertex presentation.
4. Primigravida and multipara
5. Spontaneous and induced onset of labour.

The exclusion criteria:
1. Placenta Previa.
2. Placental Abruption.
3. Previous history of postpartum haemorrhage.
4. Hemoglobin less than 7g/dL.
5. Known coagulation disorders.
6. Temperature higher than 38°C during labor (at two consecutive readings).
7. Prolonged Labor (> 20 h).
8. Accelerated Labor (< 3 h).
10. Multiple Gestations
11. Known uterine anomalies.
12. Abnormal placental adhesion
13. Forceps or Vacuum delivery.
15. History of Cesarean delivery or any uterine scar.

Ethical issues:
The hospital ethics committee approved the study.

Consent Process:
The population sample under study was instructed about research protocol and informed consent is granted from each participant before randomization.

Data analysis:
Data was collected, tabulated, then analyzed using IBM® SPSS® Statistics version 22 (IBM® Corp., Armonk, NY). Normally distributed numerical data was presented as mean and SD, and skewed data as median and interquartile range. Qualitative data was presented as number and percentage. Comparison of normally distributed numerical data was done using the unpaired Student t test. Skewed data was compared using the Mann-Whitney U test. Categorical data were compared using the chi-squared test, or Fisher’s exact test when appropriate. A two-sided p-value <0.05 were considered statistically significant.

Results

Table1: Obstetric history of patients in both study groups (age):

<table>
<thead>
<tr>
<th>Variable</th>
<th>Study group (n=75)</th>
<th>Control group (n=75)</th>
<th>t</th>
<th>Df</th>
<th>p-value¶</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>27 ± 5</td>
<td>28 ± 6</td>
<td>-0.868</td>
<td>148</td>
<td>0.387</td>
</tr>
</tbody>
</table>

Table2: Obstetric history of patients in both study groups (parity):

<table>
<thead>
<tr>
<th>Variable</th>
<th>Study group (n=75)</th>
<th>Control group (n=75)</th>
<th>χ²</th>
<th>Df</th>
<th>p-value¶</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parity PG</td>
<td>33</td>
<td>27</td>
<td>1.249</td>
<td>1</td>
<td>0.264</td>
</tr>
<tr>
<td>P1</td>
<td>15</td>
<td>16</td>
<td>21.3%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P2</td>
<td>14</td>
<td>12</td>
<td>16.0%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P3</td>
<td>5</td>
<td>11</td>
<td>14.7%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P4</td>
<td>8</td>
<td>9</td>
<td>12.0%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥P5</td>
<td>0</td>
<td>0</td>
<td>0.0%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Previous abortions Nil</td>
<td>64</td>
<td>66</td>
<td>0.372</td>
<td>1</td>
<td>0.542</td>
</tr>
<tr>
<td>One</td>
<td>8</td>
<td>6</td>
<td>8.0%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Two</td>
<td>1</td>
<td>3</td>
<td>4.0%</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Three</td>
<td>2</td>
<td>0</td>
<td>0.0%</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

The two tables show that there was no statistically significant difference detected between the 2 groups concerning, age and parity (p>0.05).
Table 3: Time of third stage took in mins. in each group.

<table>
<thead>
<tr>
<th>Group</th>
<th>Mean time to spontaneous placental separation (min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study group (n=75)</td>
<td>2.39</td>
</tr>
<tr>
<td>Control group (n=75)</td>
<td>1.93</td>
</tr>
<tr>
<td>Overall (n=150)</td>
<td>2.16</td>
</tr>
</tbody>
</table>

Log-rank test

<p>| | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>$\chi^2$</td>
<td>1.803</td>
</tr>
<tr>
<td>$Df$</td>
<td>1</td>
</tr>
<tr>
<td>$p$-value</td>
<td>0.179</td>
</tr>
</tbody>
</table>

This table shows the time of third stage took in seconds in each group with no statistically significant difference detected ($p>0.05$).

Fig. 1: Kaplan-Meier curves for time to spontaneous placental separation in both study groups.

Fig. 2: Mean time to spontaneous placental separation. Error bars represent the standard error of the mean.

Table 5: Blood lost in ml in third stage of labour in each group.
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<table>
<thead>
<tr>
<th>Variable</th>
<th>Study group (n=75)</th>
<th>Control group (n=75)</th>
<th>U</th>
<th>Z</th>
<th>p-value¶</th>
</tr>
</thead>
<tbody>
<tr>
<td>Estimated blood loss (ml)</td>
<td>200 (100 – 300)</td>
<td>200 (100 – 300)</td>
<td>2793.0</td>
<td>-0.074</td>
<td>0.941</td>
</tr>
</tbody>
</table>

This table shows the amount of blood loss in ml in third stage of labour in each group with no statistically significant difference detected (p>0.05).

![Box plot showing blood loss in ml in third stage of labour in each group.](image)

Fig. 3: Blood lost in ml in third stage of labour in each group.

Table 6: Hemoglobin pre-partum in gm/dl in each group.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Study group (n=75)</th>
<th>Control group (n=75)</th>
<th>T</th>
<th>df</th>
<th>p-value¶</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-delivery hemoglobin (g/dl)</td>
<td>10.7 ± 1.0</td>
<td>10.7 ± 0.9</td>
<td>-0.307</td>
<td>148</td>
<td>0.759</td>
</tr>
</tbody>
</table>

This table shows the hemoglobin pre-partum in gm/dl in third stage of labor in each group with no statistically significant difference detected (p>0.05).

Table 7: Hemoglobin postpartum in gm/dl in each group.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Study group (n=75)</th>
<th>Control group (n=75)</th>
<th>t</th>
<th>Df</th>
<th>p-value¶</th>
</tr>
</thead>
<tbody>
<tr>
<td>Post-delivery hemoglobin (g/dl)</td>
<td>10.4 ± 1.0</td>
<td>10.5 ± 0.9</td>
<td>0.705</td>
<td>148</td>
<td>0.482</td>
</tr>
</tbody>
</table>

This table shows the hemoglobin postpartum in gm/dl in third stage of labor in each group with no statistically significant difference detected (p>0.05).

Table 8: Drop in hemoglobin in both study groups.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Study group (n=75)</th>
<th>Control group (n=75)</th>
<th>U</th>
<th>Z</th>
<th>p-value¶</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drop in hemoglobin (g/dl)</td>
<td>0.3 (0.2 – 0.4)</td>
<td>0.2 (0.2 – 0.3)</td>
<td>2431</td>
<td>1.47</td>
<td>0.142</td>
</tr>
<tr>
<td>Percentage of drop in hemoglobin</td>
<td>2.7 (1.8 – 3.7)</td>
<td>2.1 (1.8 – 3.2%)</td>
<td>2415</td>
<td>1.494</td>
<td>0.1351</td>
</tr>
</tbody>
</table>

This table shows the drop in hemoglobin in gm/dl in third stage of labor in each group with no statistically significant difference detected (p>0.05).

Discussion
In our study, there was no statistically significant difference between the two groups according to maternal age and parity. In the group A: (the experimental group: 75 women) ≥10 IU (1mL) oxytocin was injected into the umbilical vein at the most proximal site to the placenta after clamping and cutting of the umbilical cord. In the group B: (control group: 75 women) 1 mL normal saline was injected into the umbilical vein at the most proximal site to the placenta after clamping and cutting of the umbilical cord, this is in agreement with Ghulmiyyah et al. (6) who used the same drugs which used in this study but on number of the patients more than that in the present study.

Also in our study, there was no statistically significant difference between the two groups regarding to time of third stage took in minutes with mean 2.39 min in (study group),
1.93 min in (control group). P value 0.179. This goes with the results of study carried out by Ghulmiyyah et al. (6) who found mean time of third stage took in minutes 4.1 min (Study group), 3.2 min. They believe that this lack of difference may be the result of a type II error; the difference in the mean duration of the third stage of labor between the two groups was only 2 minutes, and their sample size was inadequate to show a difference in this outcome.

In Sharma et al. (7) randomly assigned 958 women into 2 groups. Group I placental cord drainage and group II administration of with delivery of the anterior shoulder or immediately after cord clamping. The third stage had a mean duration of 3.2410 minute in oxytocin intraumbilical injection group and 3.20 minutes in cord drainage group with p value 0.157. So there was no statistically significant difference between the two groups according to the duration of third stage of labor.

In Ojha and Malla (8) study, there was no difference in the duration of third stage of labour (3.6 vs. 3.7min) between the intraumbilical and intramuscular groups.

In our study, there was no statistically significant difference regarding blood loss in the third stage of labor with Mean blood loss was 200 ml in both group, this was in accordance with Dickinson et al. (9).

Conclusion and Recommendations

Intra-umbilical oxytocin is a useful alternative in patients where methylergometrine is contraindicated or in cases where intravenous fluids need to be restricted. For optimum effect, rapid injection immediately after clamping of the cord is essential. Hence patients requiring cord blood collection, cord segment for blood gases etc, involved a time lapse and were not included in our study. Primigravidas and multigravidas requiring episiotomy showed fluctuations in the results due to variations in the blood loss. However, intra-umbilical injections can be used in both these groups.

References