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Active management of the third stage of labour with and without controlled cord traction: a randomised, controlled, non-inferiority trial

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Summary

Background Active management of the third stage of labour reduces the risk of post-partum haemorrhage. We aimed to assess whether controlled cord traction can be omitted from active management of this stage without increasing the risk of severe haemorrhage.

Methods We did a multicentre, non-inferiority, randomised controlled trial in 16 hospitals and two primary healthcare centres in Argentina, Egypt, India, Kenya, the Philippines, South Africa, Thailand, and Uganda. Women expecting to deliver singleton babies vaginally (ie, not planned caesarean section) were randomly assigned (in a 1:1 ratio) with a centrally generated allocation sequence, stratified by country, to placental delivery with gravity and maternal effort (simplified package) or controlled cord traction applied immediately after uterine contraction and cord clamping (full package). After randomisation, allocation could not be concealed from investigators, participants, or assessors. Oxytocin 10 IU was administered immediately after birth with cord clamping after 1–3 min. Uterine massage was done after placental delivery according to local policy. The primary (non-inferiority) outcome was blood loss of 1000 mL or more (severe haemorrhage). The non-inferiority margin for the risk ratio was 1·3. Analysis was by modified intention-to-treat, excluding women who had emergency caesarean sections. This trial is registered with the Australian and New Zealand Clinical Trials Registry, ACTRN 12608000434392.

Findings Between June 1, 2009, and Oct 30, 2010, 12,227 women were randomly assigned to the simplified package group and 12,163 to the full package group. After exclusion of women who had emergency caesarean sections, 11,861 were in the simplified package group and 11,820 were in the full package group. The primary outcome of blood loss of 1000 mL or more had a risk ratio of 1·09 (95% CI 0·91–1·31) and the upper 95% CI limit crossed the pre-stated non-inferiority margin. One case of uterine inversion occurred in the full package group. Other adverse events were haemorrhage-related.

Interpretation Although the hypothesis of non-inferiority was not met, omission of controlled cord traction has very little effect on the risk of severe haemorrhage. Scaling up of haemorrhage prevention programmes for non-hospital settings can safely focus on use of oxytocin.

Funding United States Agency for International Development and UN Development Programme/UN Population Fund/WHO/World Bank Special Programme of Research, Development and Research Training in Human Reproduction, Department of Reproductive Health and Research.

Introduction Post-partum haemorrhage is a major cause of severe morbidity and maternal death, particularly in Africa and Asia, where nearly a third of pregnancy-related deaths are associated with haemorrhage. Most such deaths occur because of insufficient uterine contraction soon after birth. Two management packages for the third stage of labour are commonly used, known as active management and expectant management. In active management, several prophylactic interventions are applied in combination. WHO recommends administration of oxytocin soon after delivery of the baby, controlled cord traction, and delayed clamping and cutting of the cord until the health-care worker is ready to apply traction. Uterine massage after placental delivery is included in professional society guidelines. In expectant management, the interventions included in active management are withheld unless needed. Randomised trials of active versus expectant management have been done in hospital settings and they included early clamping and cutting of the cord in addition to the WHO components. Overall, the risk of post-partum haemorrhage was more than 60% lower with active management than with expectant management. The timing of cord clamping does not seem to play a significant part in blood loss. Side-effects such as increased blood pressure, nausea, vomiting, and increased placental retention are generally attributed to the use of uterotonic ergot alkaloids.

WHO recommendations published in 2007 advocated use of the full active management package, while...
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Method

Study design and patients

We did a multicentre, non-inferiority, randomised controlled trial in 16 hospitals and two primary health-care centres in Argentina, Egypt, India, Kenya, the Philippines, South Africa, Thailand, and Uganda. Before the trial started we assessed the existing policies at the study sites. With the exception of the two primary-care centres in India, all sites practised controlled cord traction. All sites used intramuscular or intravenous oxytocin 10 IU, although the timing of administration differed. In the Philippines, a policy of concurrent use of intramuscular oxytocin and ergometrine was in place before the trial. In Argentina and Kenya, cord clamping was described as delayed, whereas other sites were practising immediate cord clamping. Uterine massage after placental delivery was part of routine management in the Philippines and Thailand and reported as about 50% in Egypt. It was not part of routine policy in any of the other countries.

All women expecting to deliver vaginally at the participating hospitals were potentially eligible. They were excluded if they were first seen in advanced labour (>6 cm cervical dilatation), were not capable of giving consent because of an obstetric emergency, a mental disorder, or distress, or if they planned to have caesarean section, were minors without a guardian, or had twin pregnancies. The fetus had to be at a gestational age of viability according to local limits. Women were asked for written informed consent either before or early in labour.

The trial protocol was approved by the WHO Ethics Review Committee and the local ethics committees of the participating institutions.

Randomisation and masking

Random allocation took place as close to the anticipated vaginal delivery as possible. The random allocation sequence was computer generated centrally at WHO. Randomisation was stratified by country and restricted with randomly varying blocks of six to eight. At each facility, a computer programme with the random allocation sequence was provided and allocation was made once the woman’s details were entered into the computer by local investigators. Each site had one spare computer in case of break-down or theft; if both failed the centre had to revert to sealed opaque envelopes as the back-up option. Centres in Egypt and Kenya had to revert to sealed opaque envelopes because of break-down and theft midway through the trial. Neither the investigators nor participants could be masked to the interventions or outcomes.

Procedures

In both groups of the trial, intramuscular oxytocin 10 IU was administered as soon as possible after birth, preferably within 1 min. If a woman had an intravenous line, oxytocin diluted in 100 mL saline could be administered through the line for 5 min. The cord was clamped and cut after manual or visual observation of a uterine contraction, usually around 1–3 min after delivery of the baby. Collection of lost blood was initiated immediately after birth of the baby by passing a drape under the woman’s buttocks. Blood collected in the drape was put in a bucket, weighed on a digital scale together with the drape, and the amount recorded in grams. Trial interventions were administered by the staff who managed the deliveries. For the analysis, the weight of the drape was subtracted and the weights were converted to volumes; the value in grams was divided by 1.06 (blood density in g/mL).

The experimental intervention assessed in the trial was the simplified package, in which placental delivery was allowed to occur with the aid of gravity and maternal effort. The birth attendant was expected to note the signs of placental separation such as a gush of blood, uterine contraction, lengthening of the umbilical cord, and visualisation of the placenta in the vagina, and then encourage the woman to cough or push. The control was the full package, in which controlled cord traction was applied immediately after observation of a uterine contraction. The cord was clamped and cut as described for the simplified package. The full package practised in the trial was applied as in other active management trials except for timing of cord clamping. However, available evidence suggests that the timing of cord clamping is not likely to have a major effect on blood loss. The protocol required that after placental delivery the uterine
fundus was rubbed and any clots expressed. Because policies for uterine massage varied and no evidence to support or refute its role is available, individual sites were allowed to implement it or not according to their existing routine. The protocol stated that, if the procedure was used, the uterus should be massaged gently until it contracted and then every 15 min for up to 2 h. Adherence to allocated treatment was defined as hands-off management of the cord in the simplified package group and application of cord traction within 30 min of delivery in the full package group.

The primary (non-inferiority) outcome was severe post-partum haemorrhage defined as measured blood loss of 1000 mL or more at 1 h and up to 2 h for women who continued to bleed after 1 h. The secondary (superiority) outcomes were blood transfusion, use of additional uterotonic, blood loss of 500 mL or more, maternal death, manual placental removal, surgical procedures (hysterectomy, ligation of vessels), the composite outcome of maternal death or severe morbidity (admission to intensive-care unit, hysterectomy, blood loss of 2000 mL or more, uterine inversion), and initiation of breastfeeding.

The Data Safety and Monitoring Committee met three times during the course of the project. The Committee reviewed two interim analyses according to the Haybittle-Peto rule with a superiority criterion on the primary outcome. The committee advised the trial Steering Committee to continue recruitment on both occasions.

Statistical analysis
We aimed to assess non-inferiority of the simplified package compared with the full package in terms of efficacy within a pre-stated non-inferiority margin. We chose the non-inferiority margin by examining the effect of the gold standard, in this case full active management, compared with expectant management (control) in previous trials. Estimates of severe post-partum haemorrhage with active and expectant management were taken from published and unpublished data of WHO studies in which post-partum blood loss was measured. On the basis of those data, we assumed a 1·5% risk of severe haemorrhage with the full package. The severe haemorrhage rate with expectant management was based on data available from earlier trials. Values between 3-0% and 4-0% were judged realistic. For sample size estimates, we therefore assumed a 3% risk of severe haemorrhage with the full package. The simplified package was assumed to preserve 70% of the benefit of the full active management package. With these assumptions, a trial of 22 908 women would have 80% power to show non-inferiority of the simplified package within 0·45% of the severe haemorrhage rate for the full package (ie, [1·0–0·7] × [3·0–1·5]), with a two-sided 95% CI, and an α of 2·5%. In relative terms, this gives a margin of non-inferiority of 1·3 [ie (1·5 + 0·45)/1·5 = 1·95/1·5]. The non-inferiority hypothesis was used for the primary outcome. For the secondary endpoints superiority hypotheses were used. SAS version 9.1 was used for all analyses.

We identified two analysis populations: intention-to-treat population—all enrolled women assessed within groups as randomised, irrespective of adherence—and modified intention-to-treat population—as intention-to-treat population, but excluding women having a caesarean section after randomisation. Our primary analysis and interpretation are based on the modified intention-to-treat population. We chose to focus on this population because for women having a caesarean section (whether elective or emergency), the trial interventions could not be implemented, nor the primary outcome assessed, in the same way as with a vaginal delivery. Elective caesarean section was an exclusion criterion but some emergency caesarean sections occurred after randomisation.

Although some researchers have suggested that in non-inferiority trials per-protocol analysis provides a more conservative interpretation, this notion has been disputed. In this trial, women with much blood loss in the simplified group were more likely to receive cord traction and other interventions to remove the placenta than were women with little blood loss. Because women with great blood loss would be excluded in a per-protocol analysis, the results would favour the simplified package group, and push the results towards erroneous non-inferiority.

Because the effects of uterine massage with or without the full package were unknown, we planned stratified analysis according to the policy of use or non-use of uterine massage. Finally, we did sensitivity analyses with and without the data set from the Philippines. We
Role of the funding source

The study was supported by the United States Agency for International Development and UN Development Programme/UN Population Fund/WHO/World Bank Special Programme of Research, Development and Research Training in Human Reproduction, Department of Reproductive Health and Research, WHO. The sponsors had no role in data collection, analysis, or interpretation of the data, the writing of the report, or the decision to submit for publication. All authors had access to the analysis plan, the outputs of that analysis and could see the data if they wished to do so. All authors participated in the final discussion and approved the submission.

Results

The figure shows the trial profile. Enrolment took place between June 1, 2009, and Oct 30, 2010. 3455 women were in Argentina, 3527 in Egypt, 2165 in India, 4005 in Kenya, 5051 in the Philippines, 1277 in South Africa, 3181 in Thailand, and 1729 in Uganda. Table 1 shows patient characteristics at trial entry and delivery. Adherence to the oxytocin and controlled cord traction components was high (>90%), with overall package adherence reduced mainly because of the timing of cord clamping and cutting (table 2). Rates of early cord clamping were particularly high in some of the Thai facilities because of hospital policies. Most sites did not practise uterine massage regularly, except Egypt (2507 of 3518 cases, 99.7%), Thailand (2790 of 2976, 93.7%), and Uganda (1715 of 1723 cases, 99.8%). In all other sites uterine massage was used in less than 10% of cases. Some differences between policies before and at the end of the trial are to be expected because the principal investigators attempted to bring their policies in line with trial procedures. About 6% of all women allocated to the simplified package eventually needed controlled cord traction (table 2). We were not able to differentiate whether their placentas were truly retained or whether the total included women whose placentas were already separated but not expelled by maternal efforts.

Table 2 shows primary and secondary outcomes. For the primary endpoint, the upper range of the 95% CI lies slightly above the non-inferiority limit (1·30). The risk of the secondary outcome of post-partum haemorrhage was higher in women given the simplified package than in those given the full package (risk ratio 1·07, 95% CI 1·00–1·14, superiority hypothesis). The rate of manual placental removal was significantly higher in the simplified package group (table 3). Mean blood loss was about 11 mL greater and the third stage about 7 min longer with the simplified package than with the full package. One case of uterine inversion occurred in the full package group. Other adverse events were haemorrhage-related.

In the sensitivity analysis excluding the Philippines (table 4), the summary estimates had slightly larger CIs, and the statistically significant difference in the rate of...
manual placental removal was no longer apparent. Mean blood loss was still greater and the third stage longer with the simplified package than with the full package. We recorded uterine massage after placental delivery and did a pre-specified subgroup analysis of the primary and secondary outcomes with or without uterine massage (data not shown). The interaction test did not suggest that the outcomes differed according to this policy (p=0.61).

Discussion
Our findings suggest that omission of cord traction results in very little, if any, increased risk of severe haemorrhage. The primary outcome of blood loss of 1000 mL or more has a risk ratio of 1.09 (95% CI 0.91 to 1.31) and the upper 95% CI limit crosses the pre-stated non-inferiority margin of 1.30 (appendix). If the upper limit had been below 1.30 we would have declared the simplified package non-inferior to the full package.19 The estimated number needed to harm (the reciprocal of the risk difference 0.17%) is 581 (one-sided 97.5% CI 189 to ∞), indicating that for every 581 women receiving the simplified package, one additional woman would have a severe haemorrhage than if all received the full package. We draw two inferences from these results: first, controlled cord traction is safe and its use can be continued in settings in which it is routinely practised; and second, the main component of active management is the uterotonic and in settings in which the full package cannot be used safely, focus should be on the uterotonic component (panel).

Table 3: Trial outcomes for the modified intention-to-treat population

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Simplified package</th>
<th>Full package</th>
<th>% risk difference (95% CI)</th>
<th>Risk ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blood loss ≥1000 mL*</td>
<td>239 (2%)</td>
<td>219 (2%)</td>
<td>0.17 (0.19 to 0.53)</td>
<td>1.09 (0.91 to 1.31)</td>
</tr>
<tr>
<td>Blood loss ≥500 mL*</td>
<td>1538 (14%)</td>
<td>1493 (13%)</td>
<td>0.90 (0.03 to 1.78)</td>
<td>1.07 (1.00 to 1.14)</td>
</tr>
<tr>
<td>Blood loss in mL†</td>
<td>282</td>
<td>271</td>
<td>10.8 (4.7 to 16.9)</td>
<td>−</td>
</tr>
<tr>
<td>Additional uterotonics‡‡</td>
<td>2434 (21%)</td>
<td>2390 (20%)</td>
<td>0.34 (0.69 to 1.37)</td>
<td>1.02 (0.97 to 1.07)</td>
</tr>
<tr>
<td>Blood transfusion§§</td>
<td>62 (1%)</td>
<td>55 (1%)</td>
<td>0.06 (0.12 to 0.24)</td>
<td>1.12 (0.78 to 1.62)</td>
</tr>
<tr>
<td>Manual removal of placenta¶¶</td>
<td>153 (1%)</td>
<td>105 (1%)</td>
<td>0.40 (0.14 to 0.67)</td>
<td>1.45 (1.14 to 1.86)</td>
</tr>
<tr>
<td>Third stage duration in min</td>
<td></td>
<td></td>
<td>12.6</td>
<td>6.1</td>
</tr>
<tr>
<td>Maternal death∗∗</td>
<td>2 (1%)</td>
<td>1 (1%)</td>
<td>0.01 (0.02 to 0.04)</td>
<td>2.00 (0.18 to 22.0)</td>
</tr>
<tr>
<td>Additional surgical procedures††</td>
<td>2 (1%)</td>
<td>9 (1%)</td>
<td>0.06 (0.11 to 0.00)</td>
<td>0.22 (0.05 to 1.03)</td>
</tr>
<tr>
<td>Maternal death or severe morbidity††</td>
<td>20 (1%)</td>
<td>31 (1%)</td>
<td>−0.09 (−0.22 to 0.03)</td>
<td>0.65 (0.37 to 1.13)</td>
</tr>
<tr>
<td>Baby put to breast within 30 min§§</td>
<td>10565 (90%)</td>
<td>10532 (90%)</td>
<td>0.14 (−0.62 to 0.91)</td>
<td>1.00 (0.99 to 1.01)</td>
</tr>
</tbody>
</table>

Table 4: Sensitivity analysis for the trial outcomes excluding Philippines sites, for the modified intention-to-treat population

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Simplified package</th>
<th>Full package</th>
<th>% risk difference (95% CI)</th>
<th>Risk ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blood loss ≥1000 mL*</td>
<td>153 (2%)</td>
<td>140 (2%)</td>
<td>0.14 (−0.21 to 0.49)</td>
<td>1.09 (0.87 to 1.37)</td>
</tr>
<tr>
<td>Blood loss ≥500 mL*</td>
<td>987 (11%)</td>
<td>927 (10%)</td>
<td>0.65 (−0.22 to 1.51)</td>
<td>1.07 (0.98 to 1.16)</td>
</tr>
<tr>
<td>Blood loss (mL)†</td>
<td>266</td>
<td>256</td>
<td>10.2 (3.9 to 16.4)</td>
<td>−</td>
</tr>
<tr>
<td>Additional uterotonics‡‡</td>
<td>1026 (11%)</td>
<td>962 (10%)</td>
<td>0.66 (−0.21 to 1.54)</td>
<td>1.07 (0.98 to 1.16)</td>
</tr>
<tr>
<td>Blood transfusion§§</td>
<td>50 (1%)</td>
<td>34 (1%)</td>
<td>0.17 (−0.02 to 0.36)</td>
<td>1.47 (0.95 to 2.27)</td>
</tr>
<tr>
<td>Manual removal of placenta¶¶</td>
<td>62 (1%)</td>
<td>64 (1%)</td>
<td>−0.02 (−0.25 to 0.21)</td>
<td>0.97 (0.68 to 1.37)</td>
</tr>
<tr>
<td>Third stage duration (min)</td>
<td></td>
<td></td>
<td>11.4</td>
<td>6.2</td>
</tr>
<tr>
<td>Maternal death∗∗</td>
<td>1 (1%)</td>
<td>1 (1%)</td>
<td>0.00 (−0.03 to 0.03)</td>
<td>1.00 (0.06 to 16.0)</td>
</tr>
<tr>
<td>Additional surgical procedures††</td>
<td>0</td>
<td>3 (1%)</td>
<td>−0.03 (−0.07 to 0.00)</td>
<td>−</td>
</tr>
<tr>
<td>Maternal death or severe morbidity††</td>
<td>12 (1%)</td>
<td>15 (1%)</td>
<td>−0.03 (−0.14 to 0.08)</td>
<td>0.80 (0.37 to 1.71)</td>
</tr>
<tr>
<td>Baby put to breast within 30 min§§</td>
<td>8571 (91%)</td>
<td>8566 (91%)</td>
<td>−0.01 (−0.85 to 0.82)</td>
<td>1.00 (0.99 to 1.01)</td>
</tr>
</tbody>
</table>

Data are n (%) or mean (SD) unless otherwise specified. N=11 821 for both packages. †Denominator is 11 821 for both groups. ‡N=11 802 for the simplified package, and 11 783 for the full package. §N=11 814 for the simplified package, and 11 790 for the full package. ¶N=11 662 for simplified package, and 11 648 for full package. **N=11 818 for the simplified package, and 11 798 for the full package. ††N=11 814 for the simplified package, and 11 790 for the full package. §§N=11 711 for the simplified package, and 11 693 for the full package.
skilled attendants are available but women prefer a hands-off mode of placental delivery. Our findings suggest such placental delivery can be safely offered in those circumstances.

Recruitment interruptions and the high number of deliveries in the Philippines resulted in loss of about 5% of forms. However, those losses were similar in both groups of the trial. The high rates of ergometrine use in one of the Philippine hospitals was another justification for the sensitivity analysis, which supported the hypothesis that the overall increase in manual removal could be attributable to ergometrine use. However, this analysis was post hoc and the increase in manual removal could be attributable to other unknown factors.

Our findings have several implications for clinical practice. Simplification of the active management of third stage of labour is possible: an intramuscular injection of oxytocin 10 IU after delivery of the baby should be regarded as the primary intervention for prevention of haemorrhage. Injections are increasingly being used in settings in which skilled birth attendants are not available but a trained health worker is present. In such settings, oxytocin should be used as the routine uterotonic for prevention of post-partum haemorrhage even if controlled cord traction cannot be implemented. Such a policy could also result in cost savings by eliminating the need for training in cord traction skills. In settings in which skilled birth attendants are available, the full package of oxytocin and controlled cord traction should be preferred especially if the shortest possible third-stage duration is desirable. Because a few women will eventually require controlled cord traction with the simplified package and it is the first procedure to be attempted in case of retained placenta, we believe that teaching of controlled cord traction in medical and midwifery curricula should continue.

Our study likewise has implications for future research. A substantial proportion of maternal deaths from haemorrhage take place in settings in which skilled birth attendants and facility-based care are not available. Our findings strengthen the need to focus on strategies to scale up the use of oxytocin in peripheral health-care settings as the primary component of active management of the third stage of labour. In this context, the use of task-shifting strategies to expand access should be considered and assessed. Such scaling-up efforts could be aided by use of prophylactic oxytocin in a prefilled, compact, auto-disabled device system. A cluster-randomised trial to investigate the effects of this system on post-partum haemorrhage at community level in Ghana is under way (NCT01108289). In settings in which no injection is possible, the role of misoprostol should be assessed with similarly rigorous research.

We were not able to examine any possible role of uterine massage in reducing blood loss in this trial. Uterine massage remains one component of third-stage management that can be implemented without any need for technology or skilled birth attendants. However, it has
not been rigorously assessed so far.\textsuperscript{21} The variation in use of uterine massage across study sites suggests that uncertainty remains among practitioners and a thorough investigation of the preventive and therapeutic use of this intervention would be timely.

**Contributors**

The idea for the study and its design were discussed and agreed at an international meeting in 2007. AMG led the writing of the draft protocol, which was commented on and finalised by the principal investigators and other members of the Steering Committee. AMG coordinated the study with support from staff in WHO (GP, MW, JPS, SJ, and EB), drafted the analysis plan with GP and DE, wrote the draft report and together with other members of the WHO team incorporated comments and suggestions from the Steering Committee members. SL did data monitoring in WHO, Switzerland, and GP analysed the data with support from SL. Site principal investigators and co-principal investigators commented on the protocol; coordinated the study in their settings; attended Steering Committee meetings; read, commented, and approved the final analysis and the article. DA, M-ES, JT, and RD were members of the Steering Committee and contributed to the protocol, participated at the SC meetings, and contributed to the final report. FA attended Steering Committee meetings, coordinated the data monitoring team in Buenos Aires, Argentina, and contributed to the protocol and the final article. DE is the Chair of the Steering Committee, contributed to the protocol, chaired Steering Committee meetings, commented on the analysis plan, drafted the final report, and interacted with the Data Safety and Monitoring Committee.

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**Investigations**

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