


# Effect of Delayed Cord Clamping at 30 Seconds and 1 Minute on Neonatal Hematocrit in Term Cesarean Delivery: A Randomized Trial

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**Objective:** To compare the effect of delayed cord clamping at 30 seconds and 1 minute on the incidence of neonatal hematocrit, anemia, maternal and neonatal complications in term cesarean delivered neonates.

**Methods:** An opened labelled, randomized controlled trial was undertaken. The 160 healthy term cesarean-born neonates were randomly allocated to either 30 seconds or 1-minute groups of delayed cord clamping (DCC) (groups 1 and 2). Neonatal venous hematocrit (Hct) and microbilirubin (Mb) were measured at 48–72 hours after birth.

**Results:** One hundred and fifty-nine neonates completed this study. Mean neonatal hematocrit  $\pm$  standard deviation at 48–72 hours was  $49.9 \pm 6.0\%$  in group 1 and  $51.2 \pm 5.9\%$  in group 2 without a statistical difference. Neonatal anemia (Hct less than 45%) occurred in 14/79 neonates (17.7%) in group 1 and in 8/80 cases (10.0%) in group 2 without a significant difference between groups. The incidence of neonatal jaundice and polycythemia (hematocrit more than 65%) was similar between groups. There were no significant differences; in the estimated blood loss during the operation, the incidence of postpartum hemorrhage and other maternal and neonatal complications.

**Conclusion:** Neonatal hematocrit was not significantly different following DCC at 30 seconds and at 1 minute, but the incidence of neonatal anemia decreased with the longer timing of DCC. The estimated blood loss and other complications were not different between the two groups. Therefore, one minute-DCC should be considered for neonatal anemic prevention when compared with 30 seconds-DCC.

**Keywords:** delayed cord clamping, timing, hematocrit, neonatal anemia

## Introduction

Neonatal anemia is an important problem that has multiple sequelae to long-term neurology, emotion, and behavior development of newborns.<sup>1,2</sup> These symptoms and sequelae can persist for more than 10 years, even if treatment was received.<sup>3</sup> Blood in the umbilical cord can be used as an autotransfusion to the neonate. Delayed cord clamping (DCC) can protect against neonatal anemia in neonates by transferring residual blood in the placenta. This practice is easy, effective and without cost.<sup>4,5</sup>

Many research studies and recommendations have supported DCC's advantages for both vaginal and cesarean delivery.<sup>5–8</sup> The proper exact time for DCC varies between authorities; for example, the World Health Organization (2012) recommends DCC at 1–3 minute,<sup>9</sup> the American College of Obstetricians and Gynecologists

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(ACOG) recommends DCC at 30–60 seconds,<sup>4</sup> the National Institute for Health and Care Excellence (NICE) recommends DCC at 1–5 minutes or longer if the mother request.<sup>10</sup>

Cesarean delivery is different from vaginal delivery. The different timing of DCC is important for mothers and neonates. A longer time for DCC may correlate with blood loss, neonatal jaundice, and polycythemia. Similarly, a shorter time for DCC may cause neonatal anemia.<sup>5</sup> The goal of this research study was to evaluate the appropriate time of either 30 seconds or 1 minute for anemic prevention in cesarean delivered neonates.

## Materials and Methods

This study was a randomized controlled trial conducted in the Department of Obstetrics and Gynecology, Udonthani Hospital, Udonthani, Thailand. It was conducted according to the Declaration of Helsinki, and national laws and regulations about clinical studies. The study protocol was approved by the Udonthani Hospital Research Ethics Committee (number 38/2561) and was registered in the Thailand Clinical Trial Registry (TCTR 20181008006). The study's participants were 160 pregnant women who underwent cesarean delivery between October 2018 and November 2019. They were counseled and invited to participate in this study.

The inclusion criteria were singleton pregnancies, age 20 years or older, who delivered by elective cesarean section between 37 and 41 weeks gestation. The exclusion criteria were pregnancy with severe medical complication, fetal anomaly, fetal growth restriction, heavy bleeding immediately after birth, or were unwilling to participate in this study. The exclusion criteria included infants with signs of birth asphyxia according to the obstetrician's judgement, such as non-vigorous infant.

A written informed consent was obtained after the explanation of the study to the participants. Then, all participants were randomly allocated to one of the two study groups. The block randomization, by a block of 4 using computer-generated numbers and sealed opaque envelopes, was performed.

The indication for cesarean delivery was according to the hospital's protocol. A preoperative blood test was done including hematocrit (Hct) and hemoglobin concentration (Hb). All cases received a prophylactic antibiotic using either cefazolin or ceftiofloxacin.

The anesthesia technique was spinal anesthesia which was performed by an anesthesiologist in the operating

room using 0.5% bupivacaine plus 0.1–0.2 mg of morphine. Then, all participants underwent low transverse cesarean section. All patients received standard care for low transverse caesarean section in the operating room, recovery room and were transferred to the postpartum ward after 2 hours post-operation.

During the operation, after delivering the infant, the research nurse announced to the obstetrician when 30 seconds had passed in group 1 and 1 minute in group 2 according to randomized group; then, the umbilical cord was clamped. During the waiting time for DCC, the baby was placed on the maternal thigh, bleeding at the uterine incision was checked and clamped. After cord clamping, the placental was delivered, intravenous oxytocin was given, the incisional wound was sutured and routine newborn care was done.<sup>8,11</sup> Blood loss during delivery was measured using visual estimation by consensus between the anesthetic and surgical nurses.

Maternal and neonatal complications, Apgar scores and birth weight were recorded. The newborn Hct and microbilirubin (Mb) were measured using venous blood at 48–72 hours after delivery. Complete blood count was analyzed by an automated hematology analyzer (Sysmex XN-3000, Meditop Company). Mb was analyzed by an automated hematology analyzer (MB NEO-BIL Plus, Zenith Science Company). Neonatal Hct was measured using a hematocrit centrifuge and a hematocrit reader.

Neonatal anemia was defined as when neonatal Hct was less than 45%, and neonatal polycythemia was when Hct was more than 65%.<sup>7,12</sup> Clinical neonatal jaundice was defined by the pediatricians using Mb level of more than the standard curve value graph<sup>13</sup> and their clinical judgement. Anemia in the mother was defined as when maternal Hct was less than 33%.

The mean neonatal Hct and prevalence of neonatal anemia at 48 hours in group 1 and 2 were compared. Neonatal polycythemia, jaundice and other neonatal complications were compared between the two groups. Estimated maternal blood loss at delivery and maternal complication between groups were also analyzed.

## Statistical Analysis

The sample size was calculated using the formula for randomized controlled trial for binary data.<sup>14</sup> The proportion of neonatal anemia at 30 seconds of DCC was 0.15 and 0.13 difference between groups were used for calculation.<sup>12</sup> A  $\alpha$  was 0.05 and the power was 80%. The calculated sample size was 72 participants in each group.

An estimated drop-out rate of 10% was added and a total number of 160 participants, with 80 per group, was used.

The participants' characteristics are presented as number, percentage, range or mean±standard deviation. The groups were compared using a linear regression analysis for continuous variables. Logistic regression analysis was used for categorical variables. The mean difference, proportion difference with a 95% confidence interval were calculated for the magnitude of effect. Data were analyzed as per-protocol method. Statistical analysis was performed using Stata version13. A P-value <0.05 was considered statistically significant.

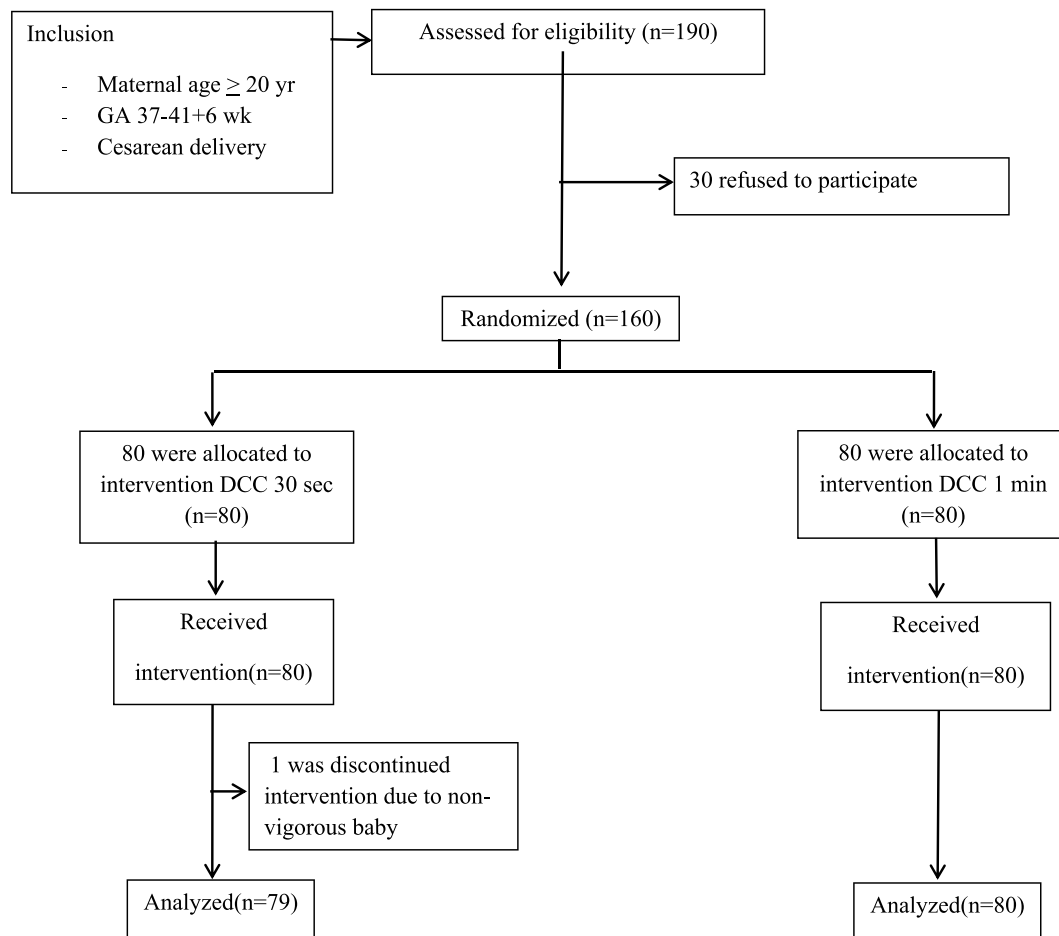
## Results

This study included 160 participants who were randomly allocated with 80 participants per group which were DCC at 30-seconds group (group1) and DCC at 1-minute group (group 2). After the study was complete, one participant in group 1 was excluded due to non-vigorous baby (no

breathing or crying). A consort diagram is presented in [Figure 1](#). The maternal baseline characteristics of each group are shown in [Table 1](#). All groups were comparable in terms of gravida, parity, gestational age, maternal age, maternal Hct and anemia, except maternal body mass index.

The mean operative time was comparable in both groups (range 23–110 minutes) and mean estimated intraoperative blood loss was 274.7 and 272.5 mL in group 1 and group 2, respectively, (range 100–800 mL) without statistically significant difference between the groups. Placenta weight and cord length were also similar in both groups.

At 48–72 hours after birth, blood sampling was done on 79 and 80 babies in groups 1 and 2, respectively. Mean venous Hct values were 49.9±6.0% in group 1 (range 37–63), and 51.2±5.9% in group 2 (range 40–65) without any statistically significant difference. Neonatal anemia (Hct less than 45%) was detected in 14/79 cases (17.7%) in group 1 and 8/80 cases (10.0%) in group 2 and there were no statistically significant differences between groups ([Table 2](#)). Neonatal polycythemia



**Figure 1** Consort diagram.

**Table 1** Comparison of Maternal Characteristics Between Groups

Characteristics	DCC 30 sec (n = 79)	DCC 1 min (n = 80)	P value <sup>β</sup>
Maternal age (yr), mean±SD	28.6±5.2	29.0±5.6	0.625
Gravida, mean±SD	2.2±1.0	2.1±0.7	0.578
Nulliparity, n (%)	23 (29.1%)	16 (20.0%)	0.299
Gestational Age (wk), mean±SD	38.9±1.1	38.8±1.2	0.304
BMI (kg/m <sup>2</sup> ), mean±SD	27.9±3.8	29.5±5.1	0.03*
Maternal Hct (%), n (%)	35.5±3.1	35.3±3.3	0.700
Anemia	6 (7.8%)	5 (6.3%)	0.738
Operation time (min), mean±SD	49.2±15.7	49.5±26.8	0.940
Cord length (cm), mean±SD	47.6±7.3	48.5±6.7	0.418
Placental weight (gm), mean±SD	551.3±102.8	569.4±109.8	0.283

**Notes:** <sup>β</sup>P value was calculated by linear regression analysis for continuous outcome and by binary regression for binary outcome. \*Statistically significant difference (P value <0.05).

**Abbreviations:** DCC, delayed cord clamping; BMI, body mass index; Hct, hematocrit; yr, year; wk, week; SD, standard deviation.

**Table 2** Comparison of Primary and Secondary Outcomes Between Groups

Characteristics	DCC 30 sec (Group 1) (n = 79)	DCC 1 min (Group 2) (n = 80)	Difference Between Group 2-1 (95% CI)	P value*
Neonatal Hct at 48-72hr (%), mean±SD	49.9±6.0	51.2±5.9	-1.3 (-3.16 to 0.56)	0.169
Neonatal anemia n (%)	14 (17.7%)	8 (10.0%)	-7.7% (-18.4 to 3.0)	0.159
Polycythemia n (%)	0 (0)	0 (0)	NA	NA
Mb at 48-72hr, mean±SD	9.4±1.9	9.2±2.3	0.13 (-0.52 to 0.79)	0.692
Clinical jaundice, n (%)	19 (24.1%)	17 (21.3%)	-2.8% (-15.8 to 10.2)	0.673
Phototherapy n (%)	15 (19.0%)	11 (13.8%)	-5.2% (-16.7 to 6.2)	0.372
Estimate blood loss (mL), mean±SD	274.7±110.3	272.5±97.1	2.2 (-30.4 to 34.7)	0.895

**Note:** \*P value was calculated by linear regression analysis for continuous outcome and by binary regression for binary outcome.

**Abbreviations:** DCC, delay clamp cord; CI, confidence interval; SD, standard deviation; Hct, hematocrit; Mb, microbilirubin.

was not found in both groups. Clinical neonatal jaundice was founded in 19 cases (24.1%) in group 1 and 17 cases (21.3%) in group 2 which was not statistically significant difference (range Mb in all cases was 3.2-18.1). Phototherapy was done in 15 cases (19.0%) in group 1 and 11 cases (13.8%) in group 2 which was similar between groups. Respiratory complications, such as respiratory distress syndrome, transient tachypnea of the newborn, and pneumonia, were similar between groups (Table 3).

The mean neonatal birthweight was 3168.8 gram (range 2250-4220) with 4 (2.5%) low birthweight babies (less than 2500 gram). The neonatal length and head circumference were also comparable between groups. Neonatal Apgar scores at 1, 5 and 10 minutes were not different between the groups (Table 3).

## Discussion

DCC has been advised by many guidelines from around the world (WHO, ACOG, and NICE)<sup>4,8,10</sup> with many studies supporting DCC's advantage in reducing anemia in

neonates.<sup>9,15-20</sup> A randomized study by Purisch et al,<sup>21</sup> compared immediate cord clamping (within 15 seconds after birth) with DCC (60 seconds after birth) in cesarean delivery. This study demonstrated that mean neonatal hemoglobin level was higher in the DCC group (18.1 g/dl vs 16.4 g/dl). However, the recommended timing of DCC in cesarean delivery is still problematic. Delayed cord clamping time in vaginal delivery can vary between 30 seconds, to more than 1 minute, or as long as the mother request but there is no definite DCC time recommended in cesarean delivery.<sup>4,8,10</sup>

During cesarean delivery, the DCC time is important for both mother and baby. Too short of a time of DCC may increase the risk of neonatal anemia, and too long of a time delays the uterine incision suturing and neonatal care. This study compared the effect of DCC at 30 seconds and 1 minute for reducing neonatal anemia. It also compared the potential for undesirable side effects of DCC such as surgical bleeding, neonatal jaundice, or polycythemia. The results demonstrate that no statistically significant difference in mean neonatal hematocrit between the groups. The incidence

**Table 3** Comparison of Neonatal Outcomes Between Groups

Characteristics	DCC 30 sec (n = 79)	DCC 1 min (n = 80)	P value*
Newborn birthweight (kg), mean±SD	3124.9±359.3	3212.7±395.9	0.146
Newborn length (cm), mean±SD	50.5±1.8	50.2±2.3	0.505
Newborn HC, mean±SD	33.5±1.2	33.6±2.9	0.709
Apgar 1 min, mean±SD	9.1±0.5	9.1±0.6	0.654
Apgar 5 min, mean±SD	10.0±0.2	9.9±0.3	0.154
Apgar 10 min, mean±SD	10.0±0.2	10.0±0.2	0.413
Respiratory complication, n (%)	3 (3.8%)	1 (1.3%)	0.305
NICU admission, n (%)	1 (1.3)	0 (0)	NA

**Notes:** \*P value was calculated by linear regression analysis for continuous outcome and by binary regression for binary outcome. Respiratory complications included respiratory distress syndrome, pneumonia and transient tachypnea of the newborn.

**Abbreviations:** SD, standard deviation; DCC, delay clamp cord; HC, head circumference; NICU, neonatal intensive care unit.

of neonatal anemia was 10.0% in 30 seconds versus 17.7% in 1 minute DCC without a significant difference.

The benefit of DCC comes from an increase in the transfusion time of residual blood in the placenta to the neonate. There are disadvantages such as increased waiting time before the cesarean incision wound is sutured, this may increase blood loss. Blood loss during cesarean delivery in this study was similar in both groups. This is also similar to Purisch et al study<sup>21</sup> that compared postoperative maternal hemoglobin between early cord clamping and DCC group. The mean neonatal blood concentration after DCC in this study was similar to a Thai report<sup>22</sup> but was lower than other western reports which the mean neonatal hematocrit of DCC infants at 24–48 hours was 55.8±5.1%<sup>7</sup> and mean hemoglobin was 18.5–18.9 g/dl.<sup>18,23</sup>

The incidence of neonatal anemia after DCC at 30 seconds in this study (17.7%) was comparable to early cord clamping in a previous study which reported 16.8% of neonates with neonatal anemia at 24–48 hours after early cord clamping.<sup>12</sup> This study's incidence of anemia in DCC at 1 minute (10.0%) was higher than a previous report which was 2.3% and 3.3% at 1 minute and 3 minutes DCC, respectively.<sup>12</sup> Performing DCC for greater than 1 minute during cesarean delivery needs additional studies to evaluate its advantage and complication.

DCC can also delay the process of newborn care including temperature care. Another worrisome complication is the increasing unnecessary blood volume to the newborn. This additional blood could lead to polycythemia, and jaundice when compared with early cord clamping.<sup>7</sup> However, this study found no significant difference in adverse neonatal effects, such as polycythemia, jaundice, or respiratory complication, in either DCC group. Therefore, our suggestion is prolonged DCC for

at least one minute during cesarean delivery which is compatible with some prior recommendations.<sup>5,8</sup>

The incidence of neonatal jaundice in this study was higher than a previous study which reported clinical jaundice in 11.5% of DCC babies and 9.9% of early cord clamping babies.<sup>7</sup> Therefore, close observation of neonatal jaundice is recommended in DCC babies. Most jaundice cases in this study were mild jaundice and all responded to the phototherapy treatment without complication.

The strength of this study is the study's design is a prospective randomized controlled trial. The limitations of this study are; first, the outcome of this study was only the short-term effect (the neonatal Hct at 48–72 hours); however, the main advantage of DCC in a term infant is the higher iron stores at 6 months of age<sup>7</sup> especially in a low iron intake area such as North-east, Thailand. Long-term study is still needed to determine the proper time of DCC effect to the infant at 6 months or older. Second, the estimation of intraoperative blood loss was done by visual estimation, so the accuracy was limited. Third, the sample size was not large enough to detect clinical assessment of safety.

## Conclusion

Neonatal hematocrit was not significantly different following DCC; at 30 seconds and at 1 minute but, the incidence of neonatal anemia decreased with the longer timing of DCC. The estimated blood loss and other complications were not different between the two groups. Therefore, one minute-DCC should be considered for neonatal anemic prevention when compared with 30 seconds DCC.

## Data Sharing Statement

All available anonymized data can be obtained by contacting the corresponding author (Dr. Metha

Songthamwat, e-mail:udonhome@yahoo.com) until 5 years after publication.

## Ethics Statement

The study protocol was approved by the Udonthani Research Ethics Committee: Number 38/2561. The study protocol was registered in the Thailand Clinical trial Registry: Number TCTR 20181008006.

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## Disclosure

The authors report no conflicts of interest in this work.

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