Predischarge transcutaneous bilirubin screening reduces readmission rate for hyperbilirubinaemia in diverse South African newborns: A randomised controlled trial

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Background. In South Africa (SA), healthy term newborns are usually discharged <72 hours after delivery. Discharged babies remain at risk for severe hyperbilirubinaemia if it is not identified early. Hyperbilirubinaemia is an important cause of readmission, and also leads to neonatal mortality and morbidity. Use of transcutaneous bilirubin (TcB) screening before hospital discharge has been controversial.

Objectives. To test the clinical benefits of TcB screening of healthy newborns before discharge for the outcomes of readmission for jaundice and severe hyperbilirubinaemia in a randomised controlled trial (RCT).

Methods. This was a RCT. We compared predischarge TcB screening with visual assessment (alone) for jaundice in apparently healthy newborns at a public tertiary hospital in Cape Town, SA. Patients or study participants were not involved in the study design and implementation.

Results. Of the 1 858 infants, 63% were black, 35% of mixed race and 1% white. There was a significant reduction in the rate of readmission for jaundice (risk ratio (RR) 0.25; 95% confidence interval (CI) 0.14 - 0.46; p<0.0001) and in the incidence of severe hyperbilirubinaemia (RR 0.27; 95% CI 0.08 - 0.97; p=0.05) with the use of TcB screening compared with visual inspection.

Conclusions. Predischarge TcB screening is superior in identifying newborns at risk of severe hyperbilirubinaemia compared with visual inspection. We recommend that every newborn, regardless of skin pigmentation, should receive objective bilirubin screening before hospital discharge. Universal bilirubin screening in newborns could potentially reduce hyperbilirubinaemia-related morbidity and mortality.

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Hyperbilirubinaemia is one of the common reasons for readmission of newborns in the first week after birth.1-3 Neonatal jaundice is usually benign and, in most instances, does not require intervention. However, up to 10% of term newborns (≥37 weeks’ gestational age) develop severe hyperbilirubinaemia that requires treatment, usually phototherapy and possibly exchange blood transfusion.4 Severe hyperbilirubinaemia is defined as a total serum bilirubin (TsB) concentration >340 µmol/L (20 mg/dL), and critical hyperbilirubinaemia as a TsB concentration >425 µmol/L (25 mg/dL) at any time during the first 28 days of life for late preterm (between 34 and 37 weeks’ gestational age) and term newborns.5 Failure to identify newborns at risk of developing severe hyperbilirubinaemia can lead to severe posticteric neurological consequences, including athetoid cerebral palsy and a variety of lifelong manifestations of kernicterus.6,7

In many settings, term newborns are discharged from hospital 1 - 2 days after delivery, usually before hyperbilirubinaemia becomes clinically evident. Peak serum bilirubin levels usually occur on postnatal days 3 - 5, when many newborns have already been discharged. The inability to identify and manage at-risk infants prior to hospital discharge in a timely manner has been cited as the major root cause of adverse outcomes of hyperbilirubinaemia.8 The burden of severe hyperbilirubinaemia continues to be high in low- and middle-income countries (LMICs). In many LMICs, severe or clinically significant neonatal hyperbilirubinaemia is not only a leading cause for hospital readmission in the first week of life, but also constitutes an important cause of neonatal mortality and morbidity.9,10 A report on the global burden of hyperbilirubinaemia suggests that sub-Saharan Africa and South Asia are the leading contributors to an estimated 1.1 million babies who develop severe hyperbilirubinaemia worldwide every year.11 In some cases, babies have levels that could lead to subsequent brain damage from kernicterus.12-14

Studies from African countries suggest that kernicterus is a leading cause of cerebral palsy15-17. In SA, a retrospective study over 5 years of exchange transfusion for hyperbilirubinaemia in newborns showed that 6 of 26 (23.0%) babies requiring exchange transfusion had signs of kernicterus.18 Findings from various observational studies conducted in North America suggest that universal bilirubin screening in newborns before discharge reduces the readmission rate for jaundice and the incidence of severe hyperbilirubinaemia.19-21 However, these findings have not been confirmed in any randomised controlled trial (RCT).22 Furthermore, there is no universal consensus on the routine screening for jaundice in all newborns using non-invasive transcutaneous bilirubin (TcB) screening or TsB measurement.23-25

The current SA guideline on the management of jaundice in the newborn does not make any recommendation on universal screening.
of all newborns with either TcB or TsB before hospital discharge.\textsuperscript{[27]} Similarly, in other African countries, there are no guidelines on universal screening of all newborns for risk of hyperbilirubinaemia. In many LMICs, newborns are assessed for jaundice by visual inspection and serum bilirubin is measured only in newborns who appear jaundiced or in some cases as part of a thyroid newborn-screening programme.\textsuperscript{[28]}

In this RCT, we compared the use of a transcutaneous device with visual inspection to screen for hyperbilirubinaemia before hospital discharge in apparently healthy term and late preterm newborns (≥35 weeks' gestational age).

We compared the impact of TcB screening before hospital discharge with visual inspection for jaundice on readmission for hyperbilirubinaemia, and the incidence of severe hyperbilirubinaemia in a SA population of newborns.

**Methods**

**Study design and setting**

This was a randomised, controlled, unblinded trial that evaluated the effect of TcB screening in term newborns before discharge from hospital. The study was conducted at the well-baby nurseries at Tygerberg Hospital, Cape Town, SA. Tygerberg Hospital is the largest tertiary healthcare institution in Western Cape Province and has 308 paediatric beds and ~7 500 deliveries per annum. According to current policy, well babies born at ≥35 weeks' gestation and with a birthweight of at least 1 800 g are discharged home after 6 hours in the case of vaginal deliveries and 48 hours in the case of caesarean sections.

**Study procedures**

We screened all infants at the time of discharge for potential eligibility into the study using the predetermined inclusion and exclusion criteria. Those who were ≤72 hours old, with a gestational age of at least 35 weeks and a birthweight of at least 1 800 g, were eligible for participation. The gestational age was determined by antenatal ultrasound scan, last menstrual period or postnatal foot length measured with a plastic Vernier's caliper for infants whose mothers were unbooked and did not have any prenatal dating scan.\textsuperscript{[29]}

**Randomisation and blinding**

Infants who met all inclusion criteria were randomly assigned to either TcB screening for hyperbilirubinaemia or visual inspection for jaundice. A random allocation sequence was generated using a computer random sequence generator. Allocation assignments were recorded on sequentially numbered sheets of paper and placed in sealed opaque envelopes, maintained in a secure location and accessible only by the principal investigator (CO). The envelopes were opened sequentially for assignment by the research assistant after informed consent was obtained and the infants allocated to the group assigned in the envelope. The study group allocation was not blinded, as it is not possible to blind clinicians or parents to the type of screening provided due to the obvious nature of the intervention. The outcome assessors and statistician were blinded to the group allocation.

**Interventions**

**TcB screening group**

In this group, we measured TcB levels once using the JM-105 Jaundice Meter (Dräger Medical, UK), a portable, handheld, non-invasive TcB measuring device, at the time of enrolment.\textsuperscript{[26]} An average of 3 readings was plotted on Bhutani's hour-specific nomogram to determine the risk zone for hyperbilirubinaemia. Infants in this group were classified into the following groups: high risk, high intermediate risk, low intermediate risk and low risk (Fig. 1).\textsuperscript{[24]} For all newborns in the high-risk zone, we obtained a TsB. The need for phototherapy was based on the TsB measurement. Infants who met the threshold for phototherapy remained in hospital for this treatment. Babies in the high-intermediate-risk, low-intermediate-risk and low-risk groups were asked to attend follow-up at a primary healthcare centre closest to their home at 24, 48 and 72 hours, respectively, after discharge home. We did not assess the timeliness of follow-up after discharge or the proportion of babies who went for follow-up.

**Visual inspection group**

Babies in this group were managed routinely according to the current standard of care. The infants were visually assessed for jaundice; the presence of tissue yellowness was assessed by blanching the skin over the glabella and sternum. The infants were evaluated by the admitting physician at the time of enrolment. Venous blood samples for TsB measurement were only collected in infants who were clinically jaundiced or in those of rhesus D-negative mothers, according to the hospital protocol. The need for phototherapy was determined based on the TsB result. Babies whose TsB value met the threshold for phototherapy according to the SA phototherapy guideline, were kept for such therapy before discharge home.\textsuperscript{[27]} Babies who did not meet the threshold for phototherapy and who were not visibly jaundiced at the time of discharge, were managed routinely. The mothers were asked to return with their newborns for follow-up assessments at the primary healthcare facility closest to their home within 2 days of hospital discharge.

**Sample size**

To the best of our knowledge, this is the first published study that assessed the rate of readmission for hyperbilirubinaemia in SA newborns. We assumed a baseline readmission rate for phototherapy of 4.8% from the literature and an unpublished local study (Ndayisaba, 2014) at Tygerberg Hospital on newborn readmission for hyperbilirubinaemia.\textsuperscript{[31–33]} We estimated

Fig. 1. Bhutani nomogram, with permission from Dr Bhutani.\textsuperscript{[24]} (TsB = total serum bilirubin.)
that a minimum of 1 858 participants would give us 80% power to detect a 50% reduction in the rate of readmission with the use of TcB screening at a 5% level of significance.

**Study outcomes**

The primary endpoint was readmission for jaundice that required phototherapy or exchange transfusion. For the primary outcome, we compared the TcB group with the visual inspection group regarding the proportion of readmissions for hyperbilirubinaemia that required phototherapy or exchange transfusion. For every infant who was readmitted, we ascertained whether they met the threshold for either phototherapy or exchange transfusion according to the SA phototherapy guideline, based on the TcB result at the time of readmission.[27]

The secondary endpoints were: phototherapy before hospital discharge, incidence of severe hyperbilirubinaemia, critical hyperbilirubinaemia or exchange transfusions and duration of hospital stay for those who were readmitted. The rate of phototherapy before discharge was measured during hospitalisation for the birth. Other outcomes (readmission for phototherapy, incidence of severe hyperbilirubinaemia and duration of hospital stay) were determined electronically through the database and also by obtaining the infants’ files and electronic patient record transcripts from the hospital or primary healthcare centre where the baby was readmitted. We obtained all TsB values during the first month of life for newborns who were readmitted. For each readmission, we confirmed the name of the hospital or primary healthcare centre where the baby was readmitted, length of stay, treatment received (phototherapy or exchange transfusion) and highest TsB level during the readmission period.

**Statistical analysis**

We collected study data using paper-based data collection forms and then transferred to the REDCap (Research Electronic Data Capture) database. Statistical analyses were done with the use of Stata 14 statistical software (StataCorp, USA).

We performed an intention-to-treat analysis. Differences according to the group allocation for continuous variables were tested with the use of unpaired t-tests with equal variances and Wilcoxon rank-sum tests. The Pearson χ² test was used to test for differences between the two groups for categorical variables. Fisher’s exact test was used to compare the rate of phototherapy before hospital discharge, readmission for hyperbilirubinaemia and incidence of severe hyperbilirubinaemia. Effect sizes were reported as risk ratios (RRs) with 95% confidence intervals (CIs). We also adjusted for any plausible baseline-confounding variable (e.g. history of jaundice in a previous sibling) by performing Mantel-Haenszel stratification.

**Patient involvement**

Patients or study participants were not involved in the design and implementation of the study.

**Ethical approval**

We obtained ethical approval from the Health Research Ethics Committee of the Faculty of Medicine and Health Sciences, Stellenbosch University (ref. no. N14/03/025). Written informed consent was obtained from the mothers for all participants before enrolment into the study. The trial was registered on clinicaltrials.gov (https://clinicaltrials.gov/ct2/show/NCT02613676).

**Results**

**Participant recruitment and follow-up**

We enrolled a cohort of 1 858 late preterm and term infants born between August 2015 and October 2016. A total of 1 910 mothers were approached for participation. Two mothers declined, and 50 were excluded because they did not meet our inclusion criteria (Fig. 2). A total of 929 infants were enrolled to each arm. All infants were screened according to the study arm to which they were allocated.

**Participant characteristics**

The baseline characteristics and demographics of enrolled infants and their mothers are shown in Table 1. There were no significant differences between the TcB screening arm and the visual inspection arm, except for the history of jaundice in a previous sibling, which was significantly higher in the TcB group.

**Outcome measures**

**Readmission for jaundice**

There was a total of 65 (3.5%) jaundice-related readmissions. We obtained the TsB results for these infants to determine if they met the threshold for phototherapy on readmission. We could not obtain the TsB results of 5 (3 in the visual inspection arm and 2 in the TcB arm) infants who were readmitted after discharge, and could not determine the appropriateness of their readmission.

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**Fig. 2. Participant study flow diagram.**
Thirteen of 929 (1.4%) infants were readmitted in the TcB arm compared with 52 of 929 (5.6%) infants in the visual inspection arm (RR 0.25; 95% CI 0.14 - 0.46; \( p < 0.0001 \)). (Number needed to treat (NNT) = 24; 95% CI 17 - 40.) Among all infants who were readmitted, there were 12 appropriate readmissions in the TcB group and 48 appropriate readmissions in the visual inspection group. As there was a randomisation imbalance in the groups regarding a previous baby with jaundice, adjustment was done by stratified analysis. The Mantel-Haenszel adjusted RR was 0.25 (95% CI 0.14 - 0.46), indicating that after adjustment for having a previous baby with jaundice, there was still a significantly protective effect of TcB screening on readmission for jaundice.

Phototherapy before discharge
The rate of phototherapy before hospital discharge was significantly higher in the TcB group than in the visual inspection group. A total of 48 (5.2%) infants received phototherapy before hospital discharge compared with 18 (1.9%) in the visual inspection group (RR 2.66; 95% CI 1.56 - 4.54; \( p < 0.0002 \)). In the TcB arm, 83 (9%) babies had predischarge TcB values in the high-risk zone (≥95th percentile); of these, 39 (47%) had TsB levels at which phototherapy was recommended according to the SA phototherapy guideline. A total of 131 (14%), 173 (19%) and 542 (58%) infants had predischarge TcB measurements in the high-intermediate, low-intermediate and low-risk zones, respectively.

Incidence of severe hyperbilirubinaemia
A total of 14 (0.75%) infants had severe hyperbilirubinaemia at the time of readmission – 3 in the TcB arm and 11 in the visual inspection arm. The rate of severe hyperbilirubinaemia was significantly lower with the use of TcB screening (RR 0.27; 95% CI 0.08 - 0.970; \( p = 0.05 \)).

Rate of exchange transfusion
Among those who were readmitted, all infants in the TcB group required phototherapy and none met the threshold for exchange transfusion. Two infants in the visual inspection group who were readmitted required exchange transfusions. This difference was not statistically significant.

Bilirubin-induced neurological dysfunction
There was 1 infant in the visual inspection group who had clinical signs that could suggest bilirubin-induced neurological dysfunction. The baby was born at 36 weeks’ gestational age with a birth weight of 2 440 g, discharged home at 72 hours after delivery, and readmitted 2 weeks later owing to prolonged neonatal jaundice. On readmission, the baby was noted to have lost about 18% of the birth weight, had a high-pitched cry and a TsB of 459 µmol/L. No magnetic resonance imaging (MRI) of the brain was done and the long-term neurological outcome for this infant is unknown. There was no documented case of infants showing neurological dysfunction in the TcB arm.

### Table 1. Baseline characteristics

| Variable                              | TcB screening (N=929), n (%) | Visual inspection (N=929), n (%) | \( p \)-value |
|---------------------------------------|------------------------------|---------------------------------|--------------|
| Gestational age (weeks), mean (SD)    | 38.4 (1.60)                  | 38.3 (1.60)                     | 0.332        |
| <37 weeks’ gestation                  | 123 (13.2)                   | 144 (15.5)                      | 0.232        |
| ≥37 weeks’ gestation                  | 806 (86.8)                   | 785 (84.5)                      | 0.212        |
| Birthweight (g), mean (SD)            | 3 128 (575)                  | 3 118 (566)                     | 0.709        |
| IUGR                                  | 106 (12.70)                  | 118 (11.41)                     | 0.393        |
| Male                                  | 485 (52.2)                   | 480 (51.67)                     | 0.816        |
| Race                                  |                              |                                 |              |
| Black                                 | 590 (63.5)                   | 585 (63.0)                      | 0.320        |
| Mixed                                 | 319 (34.3)                   | 334 (36.0)                      | 0.320        |
| White                                 | 15 (1.3)                     | 8 (0.9)                         | 0.320        |
| Other                                 | 3 (0.3)                      | 2 (0.2)                         | 0.320        |
| Breastfeeding                         | 870 (93.65)                  | 893 (96.12)                     | 0.052        |
| Cephalohaematoma                      | 1 (0.11)                     | 2 (0.22)                        | 0.056        |
| Previous sibling with jaundice        | 72 (7.75)                    | 50 (5.38)                       | 0.039        |
| Age at time of assessment, hours      | 31.1                         | 31.9                            | 0.283        |
| Mode of delivery                      |                              |                                 |              |
| NVD                                   | 492 (52.96)                  | 510 (54.90)                     | 0.287        |
| Caesarean section                     | 425 (45.75)                  | 413 (44.46)                     | 0.287        |
| Vacuum                                | 12 (1.29)                    | 6 (0.65)                        | 0.287        |
| Mother’s Rh status                    |                              |                                 |              |
| Negative                              | 16 (1.72)                    | 14 (1.51)                       | 0.927        |
| Positive                              | 871 (93.76)                  | 874 (94.08)                     | 0.927        |
| Unknown                               | 42 (4.52)                    | 41 (4.41)                       | 0.927        |
| Mother’s blood group                  |                              |                                 |              |
| A                                     | 31 (3.34)                    | 25 (2.69)                       | 0.666        |
| AB                                    | 7 (0.75)                     | 3 (0.32)                        | 0.666        |
| B                                     | 16 (1.72)                    | 16 (1.72)                       | 0.666        |
| O                                     | 37 (3.98)                    | 35 (3.77)                       | 0.666        |

TcB = transcutaneous bilirubin; SD = standard deviation; IUGR = intrauterine growth restriction (determined using the World Health Organization international growth standards for children <5 years); NVD = normal vertex delivery; Rh = rhesus.
severe hyperbilirubinaemia. [20–22,32,38,39] Our study not only validates jaundice readmissions and led to a reduction in the incidence of severe hyperbilirubinaemia in Canada have demonstrated that universal TcB or TsB reduced the incidence of the AAP and CPS guidelines, many studies in the USA and South Africa have evaluated similar outcomes. [20–22,32,38,39] We have provided evidence from a randomised controlled study. We used a proper method of randomisation and allocation concealment, with sealed opaque envelopes. Even though this method of concealment of allocation is prone to deliberate tampering, the investigators ensured that the envelopes were only opened at the time of enrolment. The study participants were similar with regard to the baseline characteristics. Although the study personnel and participants were not blinded to the treatment allocation, the outcome assessors and statistician were blinded. Therefore, we are confident that the reduction in hospital readmission rate and in the incidence of severe hyperbilirubinaemia was as a result of the TcB screening strategy in the treatment group.

Conclusions

TcB screening with risk stratification using the Bhutani nomogram in newborn infants at risk of severe hyperbilirubinaemia can help to reduce readmissions for jaundice and the incidence of severe hyperbilirubinaemia. Our study further confirms that visual inspection for jaundice is not reliable to assess the presence or absence of hyperbilirubinaemia in newborns. Universal TcB screening in all newborns before hospital discharge in SA and other African countries will be an important step towards reducing the burden and consequences of severe hyperbilirubinaemia. However, the high cost of the current TcB devices could be a barrier to implementation of universal TcB screening in many LMICs. Currently, a TcB device could cost between ZAR500 and ZAR1000. However, the use of TcB to screen for hyperbilirubinaemia in newborns has been shown to be good value for money. Therefore, nationally driven public-private solutions are warranted, such that healthcare during

Discussion

In this RCT involving indigenous South African newborns recruited from a tertiary hospital in SA, we provide evidence for the effect of predischarge TcB screening on readmission for jaundice and incidence of severe hyperbilirubinaemia. To our knowledge, this RCT is the first to evaluate the impact of TcB screening in a South African population of newborns. We demonstrated that TcB screening for hyperbilirubinaemia was superior in identifying apparently healthy newborns requiring phototherapy before hospital discharge compared with visual inspection. Earlier identification of hyperbilirubinaemia combined with treatment with phototherapy during the initial hospitalisation for the birth led to a reduction in readmission rate with the use of TcB screening. Unexpected readmission of a newborn after discharge is an indicator of inadequate assessment of the newborn’s readiness for discharge. For our study, we selected the readmission rate as a primary outcome. Although this index is often viewed from the perspective of healthcare cost, epidemiological burden and societal burden of trust between healthcare provider and the family, it can have a direct personal disruption in the maternal and newborn experience.[36–37] In 2004, the American Academy of Pediatrics (AAP) recommended a patient-friendly systems approach, i.e. that every newborn should be assessed for the risk of developing severe hyperbilirubinaemia by using predischarge TcB or TcB measurements, a policy also endorsed by the Canadian Paediatric Society (CPS).[14,15] After implementation of the AAP and CPS guidelines, many studies in the USA and Canada have demonstrated that universal TcB or TsB reduced jaundice readmissions and led to a reduction in the incidence of severe hyperbilirubinaemia.[20,22,32,38,39] Our study not only validates the finding from these studies, but also provides evidence that earlier identification of hyperbilirubinaemia with TcB screening and treatment with phototherapy is associated with a reduction in hospital readmission rates for hyperbilirubinaemia and in the incidence of severe hyperbilirubinaemia in an indigenous population of South African newborns. Other studies from LMICs have evaluated the correlation of TcB compared with TcB measurement in newborns with different skin pigmentation. These have shown a good correlation, and that implementation of TcB screening is feasible in LMICs.[14,46] However, because of varying access to healthcare and differential risk in the polycultural community of LMICs, there are limited – if any – national policy outcome studies. Therefore, we could not identify any studies from LMICs that evaluated the impact of TcB screening on readmission rate or incidence of severe hyperbilirubinaemia.
the first week after a normal pregnancy is equitable. Industry and TcB manufacturers now have the moral responsibility to make oversold transcutaneous devices safe, affordable and accessible. Reductions in the cost of readmission, the cost of more intensive treatment, and the family’s integrity are real and need to be realised for all newborns worldwide.

Declaration. This study was done as part of CO’s PhD thesis, but the publication was not a requirement for the degree.

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Author contributions. CO: conceptualised and designed the study, designed the data collection tools, co-ordinated and supervised data collection at the study site, drafted the initial manuscript and approved the final manuscript; VKB, CW, JS: provided input regarding the methods and content knowledge, critically reviewed the manuscript and approved the final manuscript; TME: conducted the statistical analysis, reviewed the manuscript and approved the final manuscript.

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Conflicts of interest. None.

Availability of data and materials. The datasets used and/or analysed during the study are available from the corresponding author on reasonable request.