A randomized control trial of phototherapy and 20% albumin versus phototherapy and saline in Kilifi, Kenya

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Abstract

Objective: The study evaluated the efficacy of phototherapy and 20% albumin infusion to reduce total serum bilirubin (TSB) in neonates with severe hyperbilirubinemia. The primary outcome was a reduction of TSB at the end of treatment. The secondary outcomes were the need for exchange transfusion, inpatient mortality, neurological outcomes at discharge, and development outcomes at 12-months follow-up.

Results: One hundred and eighteen neonates were randomly assigned to phototherapy and 20% albumin (n = 59) and phototherapy and saline (n = 69). The median age at admission was 5 (interquartile range (IQR) 3–6) days, and the median gestation was 36 (IQR 36–38) weeks. No significant differences were found in the change in TSB (Mann–Whitney U = 609, p = 0.98) and rate of change in TSB per hour after treatment (Mann–Whitney U = 540, p = 0.39) between the two groups. There were no significant differences between the two groups in the proportion of participants who required exchange transfusion (χ² (2) = 0.36, p = 0.546); repeat phototherapy (χ² (2) = 2.37, p = 0.123); and those who died (χ² (2) = 0.92, p = 0.337). Trial registration The trial was registered in the International Standardized Randomized Controlled Trial Number (ISRCTN); trial registration number ISRCTN89732754.

Keywords: Albumin, Saline, Neonates, Neonatal jaundice, Sub-Saharan Africa

Introduction

Neonatal jaundice (NNJ) is a common condition [1] and occurs when the breakdown of red blood cells is increased leading to an accumulation of bilirubin [2]. Globally, the overall incidence of NNJ is 9.9 (2.8–35.6) per 10,000 live births [1]. Africa has the highest number of children developing NNJ with an incidence of 667.8 (603.4–738.5) per 10,000 live births [1]. In Kilifi County Hospital (KCH) Kenya, NNJ accounted for 24% of young infants’ mortality and was the primary diagnosis of 17% of over 1000 neonatal admissions during the first 60 days of life in 2003 [3]. Neonates may develop hyperbilirubinemia; total serum bilirubin (TSB) above the 95th percentile for that age (in hours). Phototherapy is the treatment of choice as it enhances the conversion of bilirubin into less toxic water-soluble photo-isomers [4] and is effective in the management of NNJ [5–9]. Severe unconjugated hyperbilirubinemia can cause neurotoxicity that leads to brain damage in neonates [10] and is associated with irreversible neurodevelopmental, and neurological impairment [11–16].

The use of blue-light fluorescent tubes during phototherapy increases the need for maintenance fluid by 25% and additional intravenous fluid 1–1.5 times the maintenance fluid during intensive phototherapy [17]. The increased fluid, mainly dextrose water is needed to compensate for fluid lost in the excretion of the water-soluble photo-products from bilirubin conversion. Lack of fluid may lead to dehydration in children undergoing phototherapy [18]. Moreover, adequate hydration and urine output improve the efficacy of phototherapy.
Albumin infusion during phototherapy is reported to induce a rapid decline in unconjugated bilirubin during phototherapy [19] since albumin binds to bilirubin. Hosono et al. reported that 25% albumin infusion was more effective in reducing serum levels of unbound bilirubin during the first 6 and 24 h of treatment compared to phototherapy alone [20], but did not significantly reduce TSB levels at the end of treatment. In another study, it was reported that 10% albumin had a protective effect against bilirubin toxicity and may be useful in children with severe hyperbilirubinemia [21]. Hosono et al. further report that albumin infusion therapy is associated with better hearing outcomes at 6 months compared to standard care [22].

Despite the knowledge that albumin infusion during phototherapy may be useful, most of the evidence is from high-income countries, and less is known in sub-Saharan Africa (SSA) despite the high incidence of NNJ. The present study evaluated the efficacy of phototherapy and 20% albumin infusion versus standard care (phototherapy and saline) in the reduction of bilirubin, exchange transfusion, inpatient mortality, neurological outcomes at discharge and development outcomes at 12-months.

**Main text**

**Methods**

**Trial design**

This is a randomized controlled trial of phototherapy and 20% albumin versus phototherapy and saline. The enrolled participants were randomized in a ratio of 1:1 to the albumin group and saline group.

**Participants**

The study was conducted at KCH from 2006 to 2011. All neonates admitted to KCH had a clinical assessment of NNJ. TSB was measured in neonates (0–30 days) admitted to the KCH paediatric ward on the day of admission. Those with a TSB above 250 μmol/L were investigated to determine the cause of hyperbilirubinemia and were randomized to either phototherapy and 20% albumin or phototherapy and saline. The decision to start phototherapy was calculated based on the postnatal age, gestational, and TSB levels at admission [23]. Phototherapy was initiated when TSB levels were from 85μ/mols/L/kg in babies below 2.5 kg (as a marker of prematurity). For sick neonates, treatment was started at 30μ/mols/L below the levels for the well baby. Participants were excluded from the study if they had a congenital abnormality, clinical evidence of kernicterus, or obstructive jaundice.

**Interventions**

Neonates randomized to receive albumin were given one infusion of intravenous 20% albumin (1 g/kg) over 2 h. The neonates that were randomized to 1/5 (0.9%) normal saline sodium 154 mmol/L and chloride 154 mmol/L in 10% dextrose received the same volume of fluid as those randomized to 20% albumin i.e. 5 ml/kg over a period of 2 h. Hyperbilirubinemia was managed using Photolight, Model AS20 220–230 V with YZ20BT132/20W blue fluorescent tubes. All children were on a routine antibiotic cover (Benzyl Penicillin and Gentamicin) and maintenance fluids (10% dextrose) [23]. Discharge from the hospital occurred only when the TSB was below phototherapy level (less than 85/μmol/l/kg) and after 24 h observation to check for rebound hyperbilirubinemia after phototherapy was stopped.

Blood samples were taken for laboratory tests such as blood group determination and Coombs test (direct and indirect). Haemolytic causes were defined as mother-baby blood incompatibility or Coombs test positive. TSB was measured at admission, at 2 h, 6 h, 24 h, 48 h post admission, and every 24 h thereafter until the child was discharged. Participants were withdrawn from phototherapy if there was a need for exchange transfusion, hospital transfer, or death.

**Study Outcomes**

The primary study outcome was the reduction in TSB levels for the participants at the end of treatment. The secondary outcomes were the prevention of exchange transfusions during treatment, reduced inpatient mortality, neurological outcomes at discharge and developmental outcomes at 12-months.

**Neurological and developmental assessment**

A neurological assessments were performed using a clinical evaluation proforma designed for this study and assessed at discharge by a clinician blinded to the participants’ treatment arm. The developmental assessments were taken at 12-months of age by trained assessors. The Kilifi Development Inventory (KDI) [24, 25] and the Developmental Milestones Checklist (DMC) [24, 25] were administered to evaluate psychomotor, language and social emotional function.

**Sample size**

The sample size was based on previous data from KCH in which 48% of children with severe hyperbilirubinaemia had neurological deficits [11]. To detect a 50% difference in neurological sequelae in the albumin-treated group, assuming 80% power, 70 participants in both arms were required giving a total of 140 participants. Adjustment for death and loss to follow up of 10% required 80 participants in each arm giving a total of 160 participants.
Enrolment and randomization
The randomization was undertaken using card envelopes in blocks of 20. The random numbers were generated by the principal investigator and kept in serially numbered sealed opaque envelopes which were only opened after participants were enrolled in the study. However, due to the nature of this study, it was not possible for the investigators, assessors, clinicians, and parents to be blinded to the treatment arm once the participants were randomized.

Statistical analysis
Data were recorded on files, and double entered to a secure, Web-based REDCap database and analysed using STATA (version 15) [26]. Descriptive statistics such as frequencies, percentages, and median with respective interquartile ranges (IQR) were reported. Wilcoxon test was used to compare differences in TSB at admission and after treatment, while Mann–Whitney U test was used to compare the differences in change in TSB (the difference in TSB levels at the start and end of treatment) and rate of change per hour in TSB (the difference in the level of TSB at admission and at the end of treatment divided by the total number of hours taken on phototherapy) between the two treatments. Chi square test was used for comparisons of the proportion of participants who required exchange transfusion, repeat phototherapy, died, and participants who were discharged in the two treatments.

Results
One hundred and fifty-six participants were randomized in the study (75 assigned to 20% albumin while 76 assigned to saline).

Demographic characteristics and proportion of participants who required exchange transfusion were analysed on 118 participants (59 in the 20% albumin arm and 69 in the saline arm).

Reduction in TSB levels in the two treatments was assessed on the 70 participants (34 in 20% albumin arm and 36 in the saline arm) whose data were available (see Additional file 1: Figure S1).

Participant characteristics
The baseline characteristics of participants enrolled in the study are given in Table 1. There were no significant differences in the demographic and baseline characteristics between the participants who received phototherapy and albumin and phototherapy and saline at baseline. The median age at admission was 5 (IQR 3–6) days, the median gestation was 36 (IQR 36–38) weeks, and the median age for the mothers was 24 (IQR 19–28) years.

Most of the participants (40.8%) had haemolytic causes of jaundice, 14.6% had non-haemolytic causes, 15.4%

Table 1 Baseline characteristics participants

| Characteristics                              | 20% albumin + phototherapy (n = 59) | Saline + phototherapy (n = 69) | Test statistic |
|----------------------------------------------|------------------------------------|-------------------------------|---------------|
| Age at admission (days)                      |                                    |                               |               |
| Median (IQR)                                 | 5 (3–6)                            | 5 (4–6)                       | P = 0.574*    |
| Estimated gestational weeks                  |                                    |                               |               |
| Median (IQR)                                 | 36 (36–37)                         | 36 (36–38)                    | –             |
| Total serum bilirubin at admission           |                                    |                               |               |
| Median (IQR)                                 | 368 (303–456)                      | 392 (316–510)                 | P = 0.331*    |
| Mother's age                                 |                                    |                               |               |
| Median (IQR)                                 | 24 (19–29)                         | 24 (20–27)                    | P = 0.331*    |
| ABO incompatibility n                         |                                    |                               |               |
| Yes                                          | 28                                 | 25                            | χ²(2) = 1.85, P = 0.173 |
| No                                           | 18                                 | 28                            |               |
| Non-haemolytic causes n                      |                                    |                               |               |
| Yes                                          | 10                                 | 9                             | χ²(2) = 0.21, P = 0.646 |
| No                                           | 36                                 | 41                            |               |
| Both haemolytic and non-haemolytic n         |                                    |                               |               |
| Yes                                          | 5                                  | 3                             | χ²(2) = 0.92, P = 0.336 |
| No                                           | 54                                 | 66                            |               |
| Unknown causes                               |                                    |                               |               |
| Yes                                          | 13                                 | 18                            | χ²(2) = 0.28, P = 0.594 |
| No                                           | 46                                 | 51                            |               |

* Mann U Whitney
had both haemolytic and non-haemolytic causes, while in 29.2% of the participants the causes of jaundice were unknown.

**Reduction in bilirubin levels after treatment**
The participants’ TSB levels after treatment were significantly lower than TSB levels at admission in both the phototherapy and albumin ($Z = 5.1, p < 0.00$) and phototherapy and saline ($Z = 5.2, p < 0.001$) groups. There were no significant differences in the change in TSB (Mann–Whitney $U = 609, p = 0.98$) and rate of change in TSB per hour (Mann–Whitney $U = 540, p = 0.39$) between the two groups (Table 2).

**Outcome after treatment**
There were no significant differences ($p > 0.05$) in the proportion of participants who required exchange transfusion (three in albumin group versus four in saline group), repeat phototherapy (two in albumin versus none in saline), those who died (six in albumin group versus eleven in saline group), and participants who were discharged in the two treatments (Table 3).

**Developmental outcomes at age 12-months**
There was no significant difference in the 20% albumin group and saline group in the scores for psychomotor ($M = 26.5, SD = 11.5$, versus $M = 20.1, SD = 13.8$), $t (55) = 1.9, p = 0.060$; language ($M = 7.3, SD = 4.3$ versus $M = 5.4, SD = 4.2$), $t (55) = 1.7, p = 0.090$ and socio-emotional function ($M = 26.5, SD = 11.5$ versus $M = 25.7, SD = 11.8$), $t (55) = 0.9, p = 0.360$.

**Discussion**
This randomized control trial investigated the efficacy of 20% albumin infusion in neonates with severe hyperbilirubinemia to enhance the binding of bilirubin in the plasma and consequently reduce TSB, the need for exchange transfusion, mortality during phototherapy, neurological impairment at discharge and improve developmental outcome at 12-months. The results reveal that albumin infusion lowers the TSB levels during treatment, but there were no significant changes in TSB levels and rate of change in TSB per hour between the albumin infusion arm and the control group. Additionally, no significant differences were found in other outcomes including the need for repeat phototherapy, exchange transfusion, inpatient mortality, and developmental outcomes at 12-months.

One of the reasons for the lack of effect of albumin infusion during phototherapy could be that the extraction of bilirubin from the tissues and its discharge into the blood depend on the phototherapy rather than additional albumin infused. These findings are similar to those reported by Hosono et al. and Tsao and Victor who reported no significant reduction in TSB levels between the 25% albumin infusion group and control group. The second reason for lack of additional effect of albumin infusion during phototherapy is that albumin may lead to an increase in oncotic pressure, expanding plasma volume hence delay the excretion of bilirubin.

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**Table 2 Reduction in bilirubin levels at end of treatment in whom complete data were available**

| Variable                  | 20% albumin + phototherapy (n = 34) | Saline + phototherapy (n = 36) |
|---------------------------|-------------------------------------|-------------------------------|
| TSB at admission (SD)     | 387 (114)                           | 370 (102)                     |
| TSB after phototherapy (SD)| 191 (67)                            | 188 (69)                      |
| Change in bilirubin*      | 152 (101–248)                       | 159 (101–230)                 |
| Median (IQR)              | 2.16 (1.5–4.25)                     | 1.91 (1.19–4.00)              |

Descriptive statistics
SD standard deviation
* No significant differences in change of total serum bilirubin and rate of change in total serum reduction per hour between the two arms of the trial $p > 0.05$

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**Table 3 Outcome after treatment**

| Test statistic | 20% albumin (n = 59) | Saline (n = 69) |
|---------------|----------------------|----------------|
| Alive (%)     | 53 (89.8)            | 58 (84.1)      |
| Died (%)      | 6 (10.2)             | 11 (15.9)      |
| Repeat phototherapy (%) | 2 (3.4)               | 0 (0.0)        |
| Exchange transfusion (%) | 57 (96.6)            | 69 (100.0)     |
| Abnormal neurological functioning | –                 | –              |

$\chi^2 (2) = 0.920, P = 0.337$
from the body [20]. Third, the apparent ineffectiveness of albumin infusion may be due to oxidation of albumin during phototherapy. Oxidized albumin has a lesser affinity for bilirubin than albumin, thus promotes accumulation of free bilirubin in the body [19].

In a recent Cochrane meta-analysis evaluating the role of fluid supplementation in the treatment of NNJ, the authors reported that there were no differences between the treatment and control groups in the TSB levels 4 h after phototherapy [27]. However, several studies have reported the effectiveness of albumin infusion during exchange transfusion [21, 28, 29]. These studies have indicated that the effect of albumin infusion is useful in the clearance of bilirubin from the neonates’ tissues which may not be cleared by exchange transfusion.

Limitations
There were a number of limitations to this trial. First, the trial was not a double-blind trial, although, the endpoints of bilirubin and the need for exchange transfusions are unlikely to be biased outcomes. Second, the plasma concentrations of unbound bilirubin were not consistently measured for participants in the treatment and control group, therefore, it is difficult to determine whether the infusion of albumin affected the unbound bilirubin or not. Additionally, the developmental assessments may not have detected more subtle impairments at 12 months of age that could impact functional abilities at a later age. Finally, the study had missing data on some participants hence they could not be included in this analysis. The missing data may potentially reduce the representativeness of the sample and introduced biasness in the estimations of the outcomes of this study.

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Supplementary information
Supplementary information accompanies this paper at https://doi.org/10.1186/s13104-019-4632-2.

Additional file 1: Figure S1. Enrolment, randomization, treatment, and evaluation. The figure is a CONSORT flow diagram that shows the progress through the different phases of the parallel randomised control trial. The information provided accounts for the participants enrolled, randomized, treated, and evaluated in the study.

Abbreviations
CGMRC: Center for Geographic Medicine Research-Coast; KEMRI: Kenya Medical Research Institute; TSB: total serum bilirubin; NNJ: neonatal jaundice; KCH: Kilifi County Hospital; IQR: interquartile range.

References
1. Slusher TM, Zamora TG, Appiah D, Stanke JU, Strand MA, Lee BW, et al. Burden of severe neonatal jaundice: a systematic review and meta-analysis. BMJ Paediatr Open. 2017. https://doi.org/10.1136/bmjpo-2017-000105.
2. Ullah S, Rahman K, Hedayati M. Hyperbilirubinemia in neonates: types, causes, clinical examinations, preventive measures and treatments: a narrative review article. Iran J Public Health. 2016;45(5):558.
3. English M, Ngama M, Musumba C, Wamola B, Iwikia J, Mohammed S, et al. Causes and outcome of young infant admissions to a Kenyan district hospital. Arch Dis Child. 2003;88(S):438–43.
4. Dobbs RH, Cremer R. Phototherapy. Arch Dis Child. 1975;50(11):833.
5. Kumar P, Chawla D, Deorari A. Light-emitting diode phototherapy for unconjugated hyperbilirubinemia in neonates. Cochrane Library. 2011. https://doi.org/10.1002/14651858.CD007969.
6. Malikwe US, Jardine LA. Home-versus hospital-based phototherapy for the treatment of non-haemolytic jaundice in infants at more than 37 weeks’ gestation. Cochrane Database Syst Rev. 2014. https://doi.org/10.1002/14651858.CD007969.
7. Okwundu CI, Okoromah CA, Shah PS. Cochrane Review: Prophylactic phototherapy for preventing jaundice in preterm or low birth weight infants. Evid Based Child Health Cochrane Rev J. 2013;8(1):204–49. https://doi.org/10.1002/14651858.CD007966.pub2.
8. Sachdeva M, Murki S, Oleti TP, Kandjra H. Intermittent versus continuous phototherapy for the treatment of neonatal non-hemolytic moderate
hyperbilirubinemia in infants more than 34 weeks of gestational age: a randomized controlled trial. Eur J Pediatr. 2015;174(2):177–81. doi.org/10.1007/s00431-014-2373-8.
9. Satrom K, Slusher T, Sattom J. Effectiveness of phototherapy units in Cameroon. J Trop Pediatr. 2014;60(3):264–6. doi.org/10.1093/tropmed/ftm110.
10. Watchko JF, Tizabi C. Bilirubin-induced neurologic damage—mechanisms and management approaches. N Engl J Med. 2013;369(21):2021–30.
11. Gordon AL, English M, Tumaini Dzombo J, Karisa M, Newton CR. Neurological and developmental outcome of neonatal jaundice and sepsis in rural Kenya. Trop Med Int Health. 2005;10(11):1114–20. doi.org/10.1111/j.1365-3156.2005.01496.x.
12. Wolf M, Beunen G, Caser P, Wolf B. Extreme hyperbilirubinaemia in Zimbabwean neonates: neurodevelopmental outcome at 4 months. Eur J Pediatr. 1997;156(10):803–7.
13. Wolf M-I, Wolf B, Beunen G, Caser P. Neurodevelopmental outcome at 1 year in Zimbabwean neonates with extreme hyperbilirubinaemia. Eur J Pediatr. 1999;158(2):111–4.
14. Ogunlesi T, Dedeko I, Adekannbi A, Fetuga M, Ogundewo O. The incidence and outcome of bilirubin encephalopathy in Nigeria: a bi-centre study. Niger J Med. 2007;16(4):354–9.
15. Olusanya B, Akande A, Emokpe A, Olowe S. Infants with severe neonatal jaundice in Lagos, Nigeria occurrence, correlates and hearing screening outcomes. Trop Med Int Health. 2009;14(3):301–10. doi.org/10.1111/j.1365-3156.2009.02223.x.
16. Owa J, Davoud A. Neonatal jaundice among Nigerian preterm infants. West Afr J Med. 1990;9(4):252–7.
17. Provision Committee for Quality Improvement. Practice parameter: management of hyperbilirubinemia in the healthy term newborn. Pediatrics. 1994;94(4):558–65.
18. Iranpour R, Nohekhani R, Haghsenas I. Effect of intravenous fluid supplementation on serum bilirubin level in jaundiced healthy neonates during conventional phototherapy. J Res Med Sci. 2004;9(4):186–90.
19. Caldera R, Maynert M, Sender A, Brossard Y, Tortrat D, Galay J, et al. The effect of human albumin in association with intensive phototherapy in the management of neonatal jaundice. Archiv Francaises de Pediatrie. 1993;50(5):399–402.
20. Hosono S, Ohno T, Kimoto H, Nagoshi R, Shimizu M, Nozawa M. Effects of albumin infusion therapy on total and unbound bilirubin values in term infants with intensive phototherapy. Pediatr Int. 2001;43(1):6–11.
21. Tsao Y, Victor Y. Albumin in management of neonatal hyperbilirubinemia. Arch Dis Child. 1972;47(252):250–6.
22. Hosono S, Ohno T, Kimoto H, Nagoshi R, Shimizu M, Nozawa M, et al. Follow-up study of auditory brainstem responses in infants with high unbound bilirubin levels treated with albumin infusion therapy. Pediatr Int. 2002;44(5):488–92.
23. World Health Organization. Pocket book of hospital care for children: guidelines for the management of common childhood illnesses. Geneva: World Health Organization, 2013.
24. Abubaker A, Holding P, Van Baar A, Newton C, van de Vijver FJ. Monitoring psychomotor development in a resource-limited setting: an evaluation of the Kilifi Developmental Inventory. Ann Trop Paediatr. 2008;28(3):217–26.
25. Abubaker A, Holding P, Van de Vijver F, Bomu G, Van Baar A. Developmental monitoring using caregiver reports in a resource-limited setting: the case of Kilifi, Kenya. Acta Paediatr. 2010;99(2):291–7.
26. StataCorp L. Stata version 11.0. College Station: StataCorp LP; 2009.
27. Lai NM, Ahmad Kamar A, Choo YM, Kong JY, Ngim CF. Fluid supplementation for neonatal unconjugated hyperbilirubinemia. Cochrane Database Syst Rev. 2017;8:CD011891. doi.org/10.1002/14651858.CD011891.pub2.
28. Comley A, Wood B. Albumin administration in exchange transfusion for hyperbilirubinemia. Arch Dis Child. 1968;43(228):151.
29. Odell GB, Cohen SN, Gordes EH. Administration of albumin in the management of hyperbilirubinemia by exchange transfusions. Pediatrics. 1982;30(4):61–23.

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