

NOW GRANTED EU CONDITIONAL MARKETING AUTHORISATION.*
TECARTUS ▼ (AUTOLOGOUS ANTI-CD19-TRANSDUCED CD3+ CELLS)

IS INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MANTLE CELL LYMPHOMA (MCL) AFTER TWO OR MORE LINES OF SYSTEMIC THERAPY INCLUDING A BRUTON'S TYROSINE KINASE (BTK) INHIBITOR¹

PRESCRIBING INFORMATION

**PATIENTS WITH MCL
 POST-BTK INHIBITOR
 FAILURE FACE
 POOR PROGNOSSES²⁻⁴**

**REGAIN CONTROL
 WITH AN ORR OF
 93% WITH TECARTUS²**

(PRIMARY ENDPOINT, IN THE PRIMARY ANALYSIS SET (N=60)²)



Kaplan-Meier estimate of the duration of response, as assessed on the basis of review by the independent radiologic review committee, among 56 patients in the primary efficacy analysis who had an objective response. Tick marks indicate censored data.²
 Adapted from Wang M, et al. *N Engl J Med*. 2020.

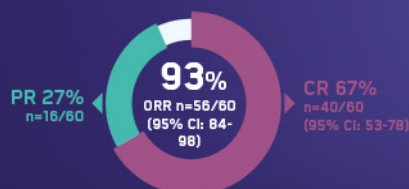
Not an actual patient.

**IN THE PRIMARY ANALYSIS SET
 (N=60) AT 12.3 MONTHS:²**

EFFECTIVE²

PRIMARY ENDPOINT:

PERCENTAGE OF PATIENTS WITH AN OBJECTIVE RESPONSE (CR OR PR)²



DURABLE

SECONDARY ENDPOINT: DOR²

The median duration of response was not reached (95% CI: 8.6-NE) at a median follow-up of 12.3 months in the primary efficacy analysis set²

- In the patients with ≥2 years follow-up, 43% (N=12/28) remained in remission²

TOLERABILITY

Tecartus led to serious and life-threatening toxic events of the type reported with other anti-CD19 CAR T-cell therapies.² The most significant and frequently occurring adverse reactions were cytokine release syndrome (91%), infections (56%) and encephalopathy (51%)¹

RAPID

Median time to response was 1 month in the primary analysis set² (range: 0.8-3.1)²

Regain control with Tecartus at www.kitecartforum.co.uk (This website contains promotional content)

ZUMA-2 was a phase 2, single-arm, open-label, multicentre trial evaluating the efficacy and safety of a single infusion of Tecartus in adult patients with R/R MCL who had previously received anthracycline or bendamustine-containing chemotherapy, an anti-CD20 antibody, and a BTKi (ibrutinib or acalabrutinib).²

*Patients are expected to enroll in a registry and will be followed in the registry in order to better understand the long-term safety and efficacy of Tecartus.¹

¹The first 60 patients treated with Tecartus who had 7 months follow-up.²

BTKi=Bruton's tyrosine kinase inhibitor; CAR=chimeric antigen receptor; CI=confidence interval; CR=complete response; DOR=duration of response; MCL=mantle cell lymphoma; NE=could not be estimated; ORR=objective response rate; OS=overall survival; PFS=progression-free survival; PR=partial response; R/R=relapsed/refractory.

REFERENCES: 1. Tecartus. Summary of Product Characteristics. 2. Wang M, Munoz J, Goy A, et al. KTE-X19 CAR T-cell therapy in relapsed or refractory mantle cell lymphoma. *N Engl J Med*. 2020;384(14):1331-1342. 3. Smith A, Roman E, Appleton S, et al. Impact of novel therapies for mantle cell lymphoma in the real world setting: a report from the UK's Haematological Malignancy Network (HMRN). *Br J Haematol*. 2018;181(215-228). 4. Smith A, Roman E, Appleton S, et al. Impact of novel therapies for mantle cell lymphoma in the real world setting: a report from the UK's Haematological Malignancy Network (HMRN) - supplementary appendix. *Br J Haematol*. 2018;181(215-228).

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TECARTUS ▼
 (autologous anti-CD19-transduced CD3+ cells)
 dispersion for infusion

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Transfusion management of severe anaemia in African children: a consensus algorithm

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Summary

The phase III Transfusion and Treatment of severe anaemia in African Children Trial (TRACT) found that conservative management of uncomplicated severe anaemia [haemoglobin (Hb) 40–60 g/l] was safe, and that transfusion volume (20 vs. 30 ml/kg whole blood equivalent) for children with severe anaemia (Hb <60 g/l) had strong but opposing effects on mortality, depending on fever status (>37.5°C). In 2020 a stakeholder meeting of paediatric and blood transfusion groups from Africa reviewed the results and additional analyses. Among all 3196 children receiving an initial transfusion there was no evidence that nutritional status, presence of shock, malaria parasite burden or sickle cell disease status influenced outcomes or modified the interaction with fever status on volume required. Fever status at the time of ordering blood was a reliable determinant of volume required for optimal outcome. Elevated heart and respiratory rates normalised irrespective of transfusion volume and without diuretics. By consensus, a transfusion management algorithm was developed, incorporating three additional measurements of Hb post-admission, alongside clinical monitoring. The proposed algorithm should help clinicians safely implement findings from TRACT. Further research should assess its implementation in routine clinical practice.

Keywords: anaemia, African children, transfusion, guidelines, malaria.

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Introduction

In sub-Saharan Africa, children admitted to hospital with severe anaemia [defined as a haemoglobin (Hb) of <60 g/l] present a major public health burden on in-patient and blood transfusion services. Although timely transfusion is frequently lifesaving, equitable access to adequate supplies of safe blood remains a key challenge.¹ In 2013 the World Health Organization (WHO) Global Database on Blood Safety reported that <5 units of blood were donated per 1000 people in most African countries,² far fewer than estimated requirements of 20 units/1000 per year.² Despite considerable investment by external funders in the decade since their previous report (2004–2005),³ few countries had made significant improvements in their collection rates.

Guidelines developed by the WHO encourage the conservative use of blood in children, restricting routine transfusion to those with severe anaemia, defined as an Hb <40 g/l, or 40–60 g/l with clinical signs of severity.⁴ However, the underpinning evidence base for this recommendation is weak⁵ and adherence to these guidelines is poor,^{6,7} which is compounded by inconsistent Hb threshold recommendations for transfusion for malaria. Currently, 20 ml/kg of whole blood (or 10 ml/kg of packed cells) is recommended for severe anaemia.⁴ However, if standard formulae for transfusion volume were applied,^{8,9} the one-size-fits-all 20 ml/kg recommendation would underestimate transfusion requirements by ~30%.^{6,10} A consensus international guideline on transfusion of critically ill children also noted a weak global evidence base and specifically highlighted the need to evaluate optimal transfusion volumes.¹¹

The Transfusion and Treatment of Severe Anaemia in African Children Trial (TRACT) was designed as a multicentre trial to establish transfusion and treatment strategies in sub-Saharan Africa and investigated: (i) whether an immediate transfusion in children with uncomplicated severe anaemia (Hb 40–60 g/l) would improve outcomes and (ii) whether 30 *versus* 20 ml/kg whole blood might improve outcomes, defined as mortality to day 28 (primary outcome) and day 180, and the need for additional transfusions and relapse of severe anaemia.¹²

The trial results, published in two linked manuscripts in 2019,^{13,14} showed that an immediate transfusion is not required in children with uncomplicated severe anaemia (Hb 40–60 g/l without severity features), as long as children are monitored for progression to the development of severe and/or complicated anaemia during their initial hospital stay; such progression occurred in 49% of trial participants who then received a transfusion.¹³ The results also showed that giving larger volumes of blood, 30 ml/kg whole blood equivalent *versus* the 20 ml/kg recommended in the current WHO guidelines, to children with severe anaemia who did not have a fever (axillary temperature $\leq 37.5^{\circ}\text{C}$, measured at screening) halved the number of deaths by 28 days [hazard ratio (HR) 0.43, 95% confidence interval (CI) 0.27–0.69].¹⁴ However, the

opposite occurred in children who were febrile ($>37.5^{\circ}\text{C}$) at screening; 30 ml/kg whole blood equivalent almost doubled the risk of death in comparison to children who received the lower, 20 ml/kg, volume (HR 1.91, 95% CI 1.04–3.49; *P* value for heterogeneity of effect on mortality *P* = 0.00009).

The TRACT results provide an evidence base with potential to improve outcomes for children with severe anaemia. Furthermore, initial economic analyses also suggested that if the results were implemented and adhered to by clinicians, there could be substantial cost savings for blood transfusion services. To explore this further, the TRACT investigators shared results from the trial with key stakeholders at a meeting in Uganda in February 2020, co-hosted by the African Society for Blood Transfusion and the Ugandan Paediatric Association. The meeting was attended by stakeholders from 12 sub-Saharan countries and included representatives from blood transfusion services, health service providers (including paediatricians involved in providing continued professional development and training courses) and representatives from international policymakers, including the WHO and Médecins Sans Frontières (Appendix 1). Our aim was to clarify current practice in different settings within Africa, to discuss the implications of the TRACT findings from a stakeholder perspective, to identify potential barriers to their uptake and to identify any additional requirements that might be needed to support their safe implementation, including further analyses of the TRACT data.

A major output from discussions at this stakeholder meeting was the development of a transfusion management algorithm for future implementation, which was agreed by consensus. Here, we present the algorithm and the additional analyses that support it.

Methods

A summary of the trial and statistical methods is available in the supplemental appendix. Blood transfusions were supplied by the local blood transfusion services (BTS), which included three blood component types: (i) whole blood (WB), collected from donors and stored without any preparation or via transfer bags into smaller bags of whole blood (known as 'Pedi-Packs'); (ii) packed red cell concentrates (RCC), produced by centrifugation to remove platelets and plasma, followed by the addition of sodium, adenine, glucose and mannitol solution; and (iii) RCC created using gravity (Uganda only).¹⁵ To support the algorithm, we investigated detailed response to transfusion volume and requirement for additional transfusions. At the request of participants at the meeting we provided original and further analyses of the primary outcome (28-day mortality) in the following key subgroups: (i) nutritional status (protocol defined); (ii) malaria status including those with high malaria burdens (plasma *Plasmodium falciparum* histidine-rich protein 2 (PfHRP2) concentration of >1000 ng/ml¹⁶); (iii) children with shock defined by the Fluid Expansion As Supportive Therapy in

critically ill African children (FEAST) trial eligibility criteria;¹⁷ and those with sickle cell disease (SCD). These subgroups all have direct implications for the current WHO management guidelines.⁴ Finally, additional exploration of the fever/no fever interaction was undertaken for a number key subgroups where the interaction might potentially have been impacted for physiological reasons [including SCD and Blackwater fever (haemoglobinuria)].¹⁸

Results

The TRACT recruited children aged 2 months to 12 years presenting to hospital with both uncomplicated and complicated severe anaemia (Fig 1), and the algorithm is aimed at this population (Fig 2). We found no evidence to show that outcome was affected by nutritional status (including children classified as undernourished) or by SCD status (which was unknown at the time of randomisation in the majority) (Fig 3A). In the current WHO guidelines, malaria hyperparasitaemia is an indicator for immediate transfusion.⁴ We therefore assessed whether this was supported by the TRACT. Instead of 'hyperparasitaemia', we used the value PfHRP2 ≥ 1000 ng/ml, a parameter that more accurately reflects the highest total parasite burden, and which distinguishes 'true' severe malarial anaemia from anaemia in the presence of incidental parasitaemia.¹⁶ We found that outcome was the same in children managed on a deferred transfusion protocol as in those receiving immediate transfusion (Fig 3A). Thus, in our algorithm we did not provide separate recommendations with regard to immediate or deferred transfusion for these groups.

For the higher *versus* lower volume comparison, the stakeholder meeting group identified two subgroups that could, potentially, have increased the risk of a worse outcome with higher transfusion volumes, irrespective of fever status: (i) children with clinical features of shock (severely impaired perfusion as defined in FEAST trial¹⁷); and (ii) children with anthropometric features of undernutrition. As shown in Fig 3B none of these subgroups had outcomes that differed significantly from the overall findings. Therefore, the stakeholder meeting concluded that the recommendations on transfusion volume did not need to differ according to a child's perfusion status, nutrition status or malaria parasite burden.

The initial evaluation of a child with suspected severe anaemia should include a Hb test, axillary (or other) temperature, weight and assessment for severity signs (altered conscious level or respiratory distress). Then the child can be grouped into one of three categories: (i) severe and complicated anaemia (Hb <40 g/l or Hb 40–60 g/l, with clinical severity features); (ii) Hb 40–60 g/l, without clinical severity features; and (iii) Hb >60 g/l, with or without severity signs. The first group require immediate transfusion; the second group do not require immediate transfusion but should have a blood sample sent to the transfusion services for blood ABO grouping¹⁹ and blood unit(s) (WB or RCC) saved while being monitored both clinically and for Hb level (at a minimum of 8, 24 and 48 h

following identification). The volume for transfusion, based on weight and temperature can be obtained from Table I. The third group does not require transfusion (with Hb >60 g/l) and can be managed without further Hb monitoring (Fig 2). Once children have been transfused, or for stable children with Hb 40–60 g/l, they will continue to be monitored and further transfusion management will depend on their latest Hb and/or clinical features.

Blood volume dosing table

In the TRACT, we were reassured that $>96\%$ of children received the correct volume of blood. Children weighed between 5 and 35 kg, the majority ($\sim 75\%$) weighing ≤ 15 kg. Thus, most of the WB prescriptions for the 20 ml/kg strategy were between 100 and 320 ml and for the 30 ml/kg strategy were between 150 and 480 ml. To simplify future implementation of the algorithm, we have generated a simplified 'look-up' chart together with bag size recommendations for WB Pedi-Packs for appropriate weight bands (Table I Blood dosing chart). If WB is unavailable then the equivalent RCC doses are half the doses stated in the 'look-up' chart. This is intended to be used alongside the transfusion algorithm.

Timing and stability of fever

For children with severe or complicated anaemia, the volume of blood component ordered (20 or 30 ml/kg WB) in the algorithm is dependent on the axillary temperature at the time of requesting blood. For those with uncomplicated anaemia that subsequently need a transfusion, the axillary temperature at the time of ordering blood would be used. In the TRACT trial nearly all children (3106/3196, 97.2%) had a history of fever in the current illness, but only 1253 (39.2%) had a recorded fever at screening, before the receipt of a transfusion. Screening temperature is only at one time point, so we examined the patterns of fever during their hospital stay and found only 10% of children without a fever at screening developed one after the transfusion had started. For children with a fever at screening, the fever had waned in 80% by 16 h (Fig 5). The heterogeneity in the estimate of mortality remained for groups defined by their fever status over 8 h (Fig 5; Figure S2) with benefit from 30 ml/kg for those with no fever at screening, even if they had intermittent fever during the subsequent 8 h. However, if a child subsequently required new or additional transfusion then the group recommended that the axillary temperature at the time of requesting blood would be used to indicate volume required, rather than the initial fever status at initial screening.

What might be the mechanisms for excess mortality in children receiving 30 ml/kg with fever?

We found no strong evidence implicating any specific cause for the excess mortality associated with fever and blood

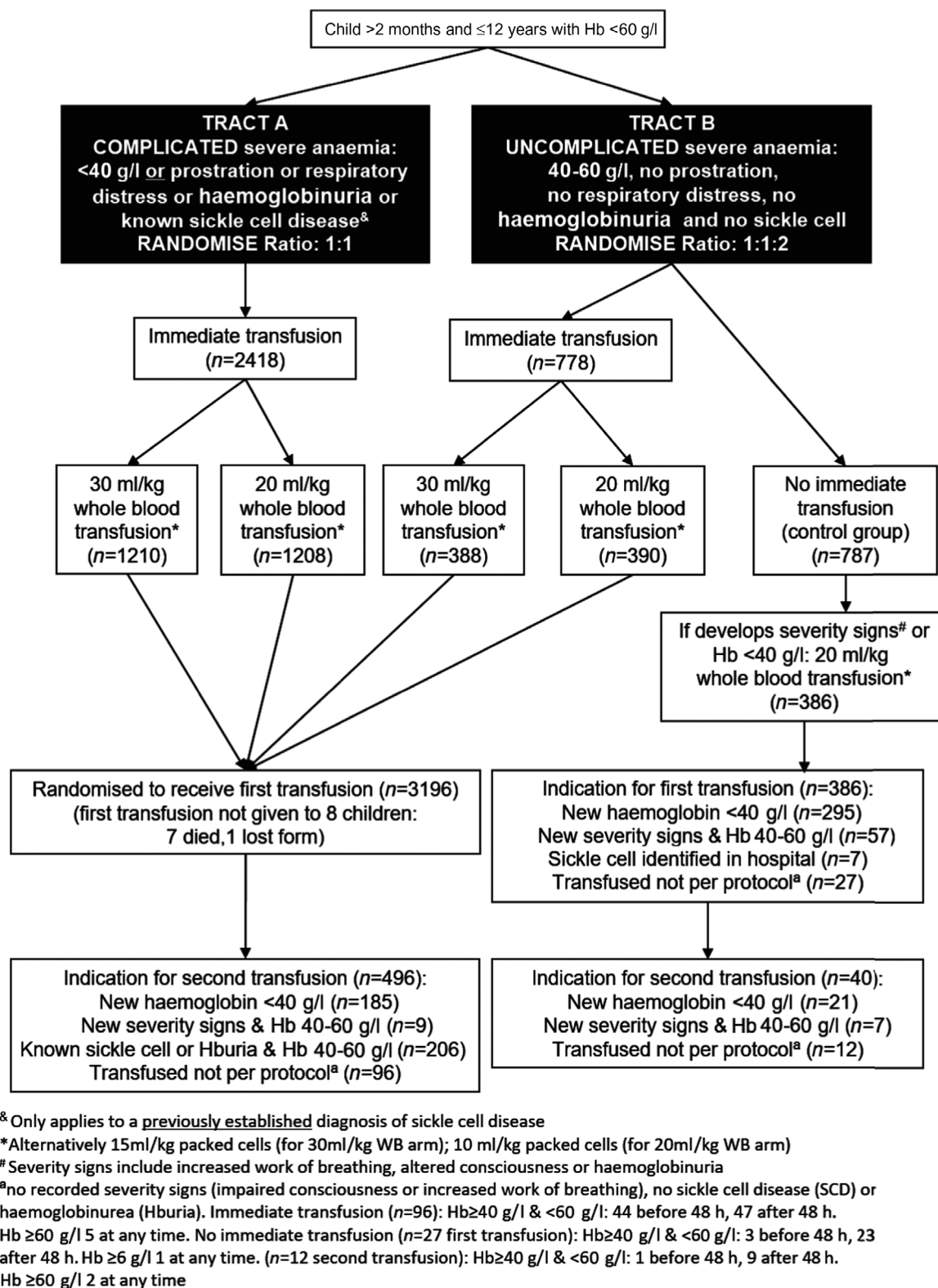


Fig 1. Transfusions given in the TRACT trial.

volume. Neither known SCD nor haemoglobinuria at admission had any effect on the fever/transfusion volume interaction, nor did the age of the child (Figure S1). Transfusion-

related volume overload [pulmonary oedema, transfusion-related lung injury (TRALI) or transfusion-associated cardiac overload (TACO)] was ruled out because of the small

Algorithm for managing suspected/confirmed severe anaemia in children aged from 2 months to 12 years

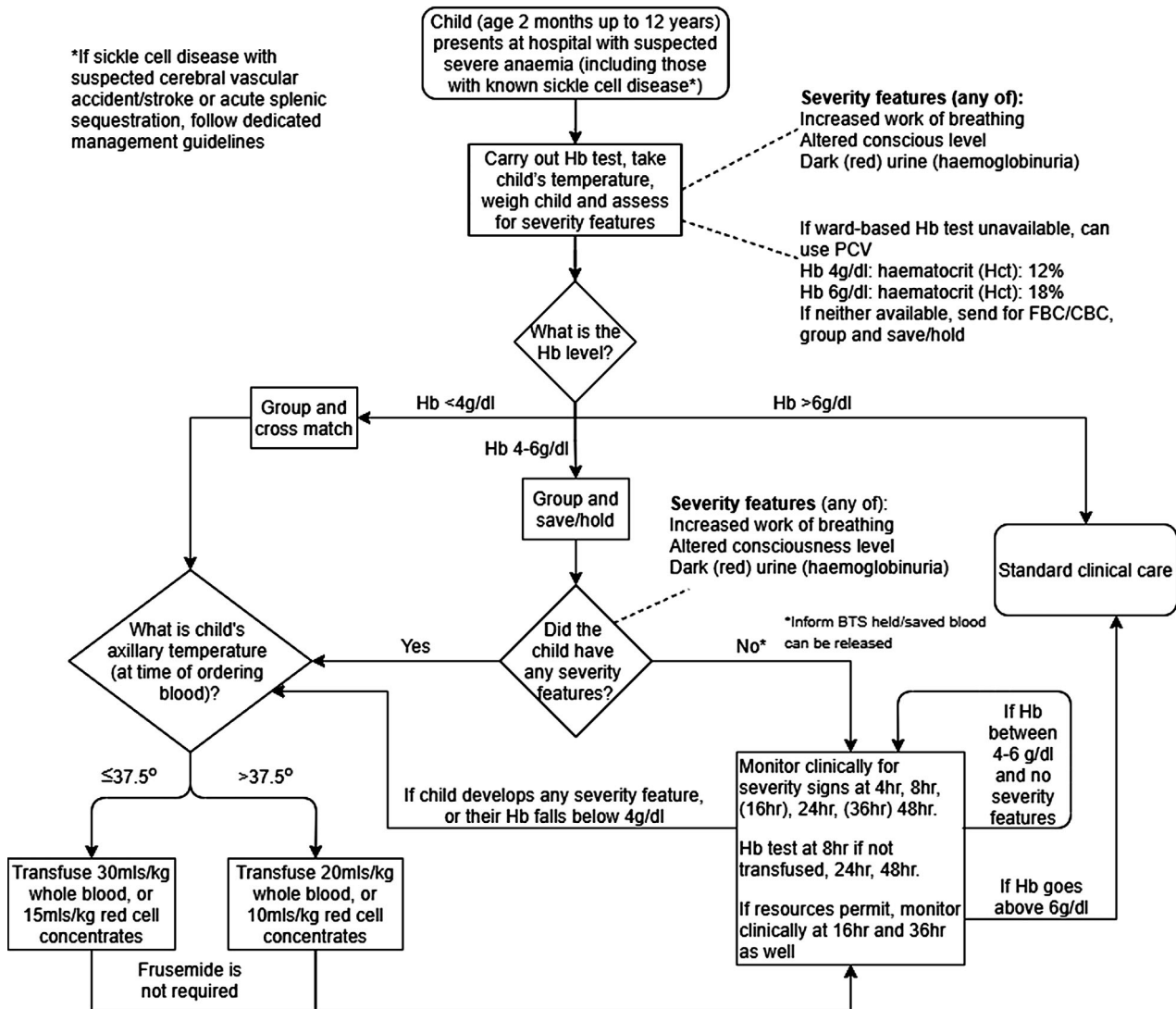
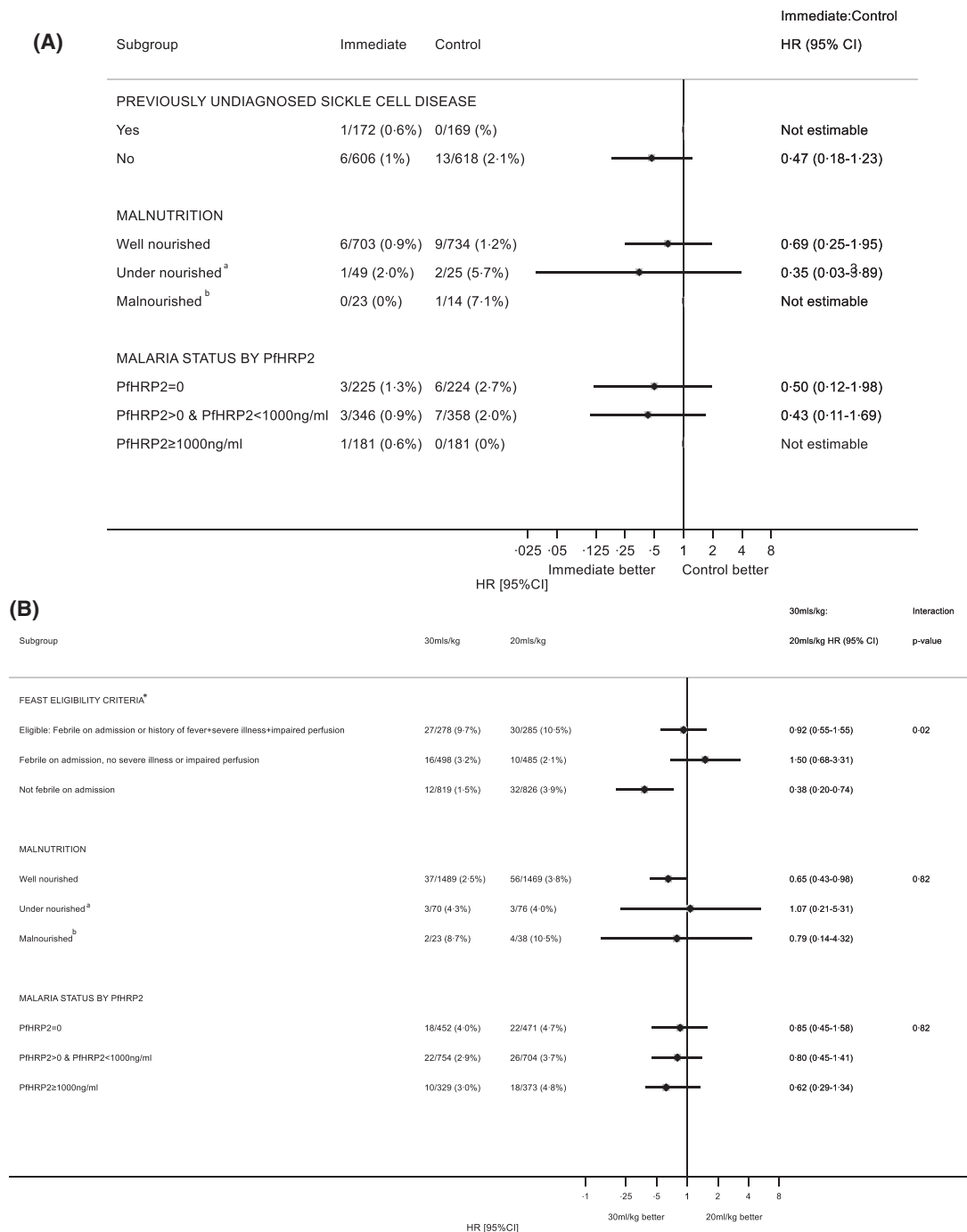


Fig 2. Algorithm for managing suspected/confirmed severe anaemia.

number of solicited events ($n = 5$) in the trial (summarised in Table S1); only one was adjudicated to be probably related to blood volume. No child received diuretics or other anti-failure medications. The recovery of elevated heart and respiratory rates occurred similarly in both febrile and afebrile groups (Figure S3). Blood component type (WB or RCC) was also explored in a pre-planned sub-analysis of the TRACT and was found not to have contributed to adverse outcomes. The stakeholders supported the algorithm recommending both 30 and 20 ml/kg transfusion WB volumes (or RCC equivalent), depending on fever status.

Children in the no immediate transfusion category: what triggered transfusion and when was it given?

Children with uncomplicated severe anaemia (no severity signs and Hb 40–60 g/l) would be monitored clinically according to the algorithm, with a Hb test at 8, 24 and 48 h and not transfused unless they develop severity signs or Hb <40 g/l. In the TRACT trial 386/787 (49%) children randomised to not receive an immediate transfusion, subsequently did receive one; details of timing of triggered transfusions are in Figs 1 and 4A. Haemodilution, by liberal use of intravenous fluids, was discounted, based on the fact



*Severe illness: respiratory distress or impaired consciousness. Impaired perfusion: one or more of the following: capillary refill >2s, temperature gradient, weak pulse or severe tachycardia. 293 children were not febrile at admission but had history of fever, severe illness and impaired perfusion and so are included in the eligible group.

^aUndernourished: one or more of as mid-upper arm circumference (MUAC) ≥ 11.0cm – 11.9cm (children aged 2 to 6 months) or MUAC ≥ 11.5cm – 12.4 (children aged 6 months to 59 months) or WHZ-3 to -2 (or WAZ if height not recorded) at any age.

^bMalnourished: one or more of mid-upper arm circumference (MUAC) < 11.0cm (children aged 2 to 6 months) or MUAC < 11.5cm (children aged 6 to 59 months) or Weight for Height Z score ((WHZ) < -3 or Weight for Age Z score (WAZ) if height not recorded) or presence of kwashiorkor at any age

Fig 3. (A) Mortality at 28 days by subgroups for the immediate vs control timing of transfusion comparison. (B) Mortality at 28 days by subgroups for the transfusion volume comparison. PfHRP2, plasma *Plasmodium falciparum* histidine-rich- protein 2.

Table I. Blood dosing chart for whole blood components.*

Weight, kg	20 ml/kg	Component†	Volume	Min, ml/kg	Mid, ml/kg	Max, ml/kg	30 ml/kg	Component†	Volume	Min, ml/kg	Mid, ml/kg	Max, ml/kg
5	6	100	100	16.7	18.2	20.0	150	150	25.0	27.3	30.0	30.0
6	7	150	150	21.4	23.1	25.0	200	2 × 100	28.6	30.8	33.3	33.3
7	8	150	150	18.8	20.0	21.4	200	2 × 100	25.0	26.7	28.6	28.6
8	9	150	150	16.7	17.6	18.8	250	250	27.8	29.4	31.3	31.3
9	10	200	2 × 100	20.0	21.1	22.2	300	2 × 150	30.0	31.6	33.3	33.3
10	11	200	2 × 100	18.2	19.0	20.0	300	2 × 150	27.3	28.6	30.0	30.0
11	12	200	2 × 100	16.7	17.4	18.2	350	250 + 100	29.2	30.4	31.8	31.8
12	13	250	250	19.2	20.0	20.8	400	150 + 250	30.8	32.0	33.3	33.3
13	14	250	250	17.9	18.5	19.2	400	150 + 250	28.6	29.6	30.8	30.8
14	15	300	2 × 150	20.0	20.7	21.4	450	Adult pack	30.0	31.0	32.1	32.1
15	16	300	2 × 150	18.8	19.4	20.0	450	Adult pack	28.1	29.0	30.0	30.0
16	17	300	2 × 150	17.6	18.2	18.8	500	Adult pack	29.4	30.3	31.3	31.3
17	18	350	250 + 100	19.4	20.0	20.6	500	Adult pack	27.8	28.6	29.4	29.4
18	19	350	250 + 100	18.4	18.9	19.4	550	Adult + 100	28.9	29.7	30.6	30.6
19	20	400	150 + 250	20.0	20.5	21.1	600	Adult + 150	30.0	30.8	31.6	31.6
20	21	400	150 + 250	19.0	19.5	20.0	600	Adult + 150	28.6	29.3	30.0	30.0
21	22	400	150 + 250	18.2	18.6	19.0	650	Adult + 150	29.5	30.2	31.0	31.0
22	23	450	Adult pack	19.6	20.0	20.5	650	Adult + 150	28.3	28.9	29.5	29.5
23	25	450	Adult pack	18.8	19.1	19.6	700	Adult + 250	29.2	29.8	30.4	30.4
>25	450	Adult pack	Adult pack	18.0	18.4	18.8	700	Adult + 250	28.0	28.6	29.2	29.2

Shaded rows are WHO standard weight bands.

*If whole blood packs are not available use red cell concentrate (RCC)/packed cells at half the volume, e.g. 20 ml/kg whole blood is equivalent to 10 ml/kg RCC.

†If multiple blood components are needed then pragmatically these can be obtained from multiple donors if a single donor component is insufficient.

that no fluid boluses were given¹⁷ and children only received maintenance fluids until they were able to drink. Compared to children randomised to immediate transfusion, who received blood at a median [interquartile range (IQR)] of 1.3 [0.9–1.7] h after randomisation, 295/386 (76%) of those in the deferred group received a subsequent transfusion, principally triggered by a fall in Hb to <40 g/l, at a median (IQR) of 24.9 (9.2–49.8) h after randomisation. Reasons for transfusion in an additional 57 (15%) children were the development of clinical severity features (impaired consciousness or increased work of breathing) and SCD identified in hospital ($n = 7$; 2%). Only 27 (7%) had a transfusion without a very low Hb or a recorded severity feature, largely after 48 h when reasons for transfusion were not routinely captured. Of note, the timing of triggered transfusions largely coincided with the routine Hb assessment in the trial (at 8, 16, 24 and 48 h after randomisation) and 49% were given within 24 h (excluding the 24 assessment) (Fig 4A). A further two Hb measurements (at 36 and 48 h) would have identified around two-thirds (249/386; 64.5%) of children in the control group who subsequently received a transfusion. Thus, this informed the recommendation in the algorithm for repeat Hb measurements at 8, 24 and 48 h (Fig 2), with additional measurements at 16 and 36 h where resources permit.

Timing of deaths in controls: concern that this may be related to delaying transfusion

Of the 13 children in the control arm who died, 10 had received a transfusion prior to death and these were received at a median of 9 h after randomisation.¹³ Furthermore, of seven children transfused before 10 h, one died on Day 1 (who had a screening Hb of 59 g/l) and six died between days 4–23, with a recorded 48-h Hb level of 50–90 g/l (only two were <60 g/l). Three were not transfused; one died on day 1 with last recorded Hb of 46 g/l, while two died on day 6 and day 25 with 48-h Hb levels of 54 and 65 g/l respectively. In summary, delay in transfusion or no receipt of transfusion does not appear to have made a clear contribution to outcome among these deaths (Supplemental material original report).¹³

In children requiring additional transfusions: what triggered re-transfusion events and when were they given?

In the TRACT trial, of those receiving immediate transfusion, 3188/3196 (99%) received one, and 496/3196 (15.5%) received a second transfusion, 300 (19%) in the 20 ml/kg group and 196 (12%) in the 30 ml/kg group, an absolute difference of 6% (95% CI 4–9%, $P < 0.0001$) greater in the 20 ml/kg strategy.¹⁴ Overall, re-transfusions were due to falls in Hb to <40 g/l in 185 (37%) children, and occurred more frequently in the 20 ml/kg group (42% vs. 30% in 30 ml/kg

group) (Table II), or it was felt that clinically stable children with SCD or haemoglobinuria with Hb between 40 and 60 g/l needed it corrected to >60 g/l ($n = 206$, 42%). The number for which an Hb <40 g/l prompted the second transfusion supports our recommendation in the algorithm for repeat Hb measurements at 8, 24 and 48 h among children who have been transfused, as well as those being monitored before a possible transfusion. These three additional measurements capture the large proportion of repeat transfusions, as very few received an additional transfusion for new clinical signs of severity after admission (2%) or when Hb was >60 g/l (Fig 1). With regard to the timing of additional transfusions, they occurred earlier and at a higher rate in those randomised to 20 ml/kg. Approximately 5% of children in this group received a transfusion by 24 h and 9–10% by 48 h, compared to under half this rate at both time points in the 30 ml/kg arm (Fig 4B). Although it was the second most common reason for a second transfusion, SCD status (known or unknown) was not a risk factor for poor outcome, this group was not segregated in the algorithm for repeat transfusion.

Consensus blood transfusion algorithm

Following discussion of all of the above, the meeting participants endorsed the proposed paediatric blood transfusion algorithm based on the evidence provided by the TRACT (Fig 2). The key elements start with triage of a child with suspected severe anaemia, incorporating an assessment of severity (to determine the need for a transfusion), with further management guided initially by an assessment of Hb (or haematocrit if Hb is not possible) and fever status at the time a blood transfusion is ordered from the blood transfusion service, in order to determine whether to request a higher volume of blood (30 ml/kg WB or 15 ml/kg RCC), or standard volume (20 ml/kg WB or 10 ml/kg RCC) if fever is present. Informed by the main time points when first (for children with uncomplicated anaemia) or repeat transfusions occurred, our proposed algorithm also incorporates a minimal number of additional measurements of Hb (or haematocrit) and clinical monitoring.

Discussion

If fully implemented, our proposed paediatric blood transfusion algorithm could avert the need for immediate transfusion in children with severe uncomplicated anaemia (including those with SCD) and provide higher volumes of blood to children with severe and complicated anaemia (based on clinical signs and severity of anaemia) who do not have a fever. Both these strategies would improve clinical outcomes, more specifically, in children without a fever, a liberal 30 ml/kg strategy would result in almost 60% fewer children dying, while for children with a fever a conservative 20 ml/kg strategy would prevent a 50% higher mortality were

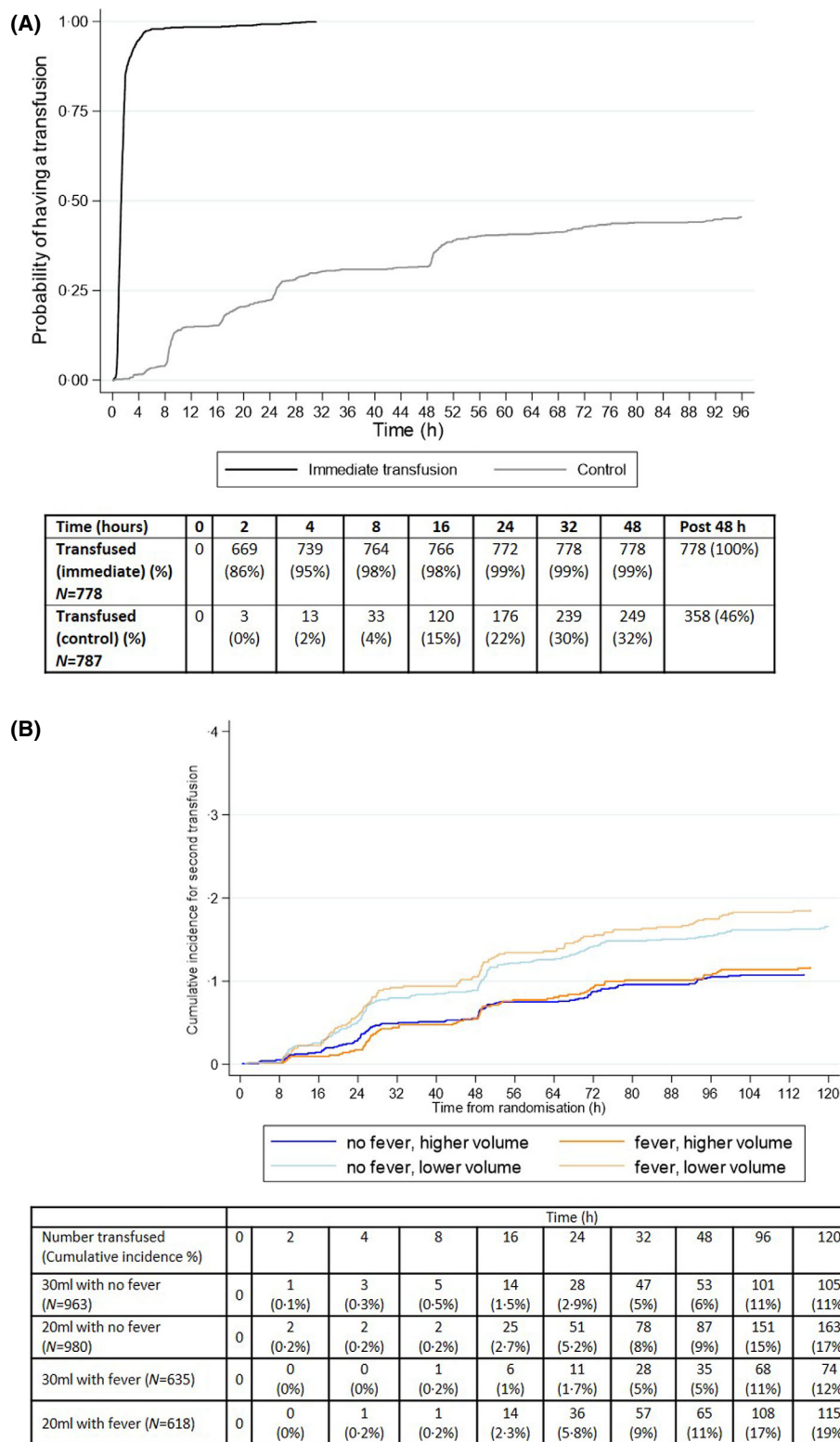
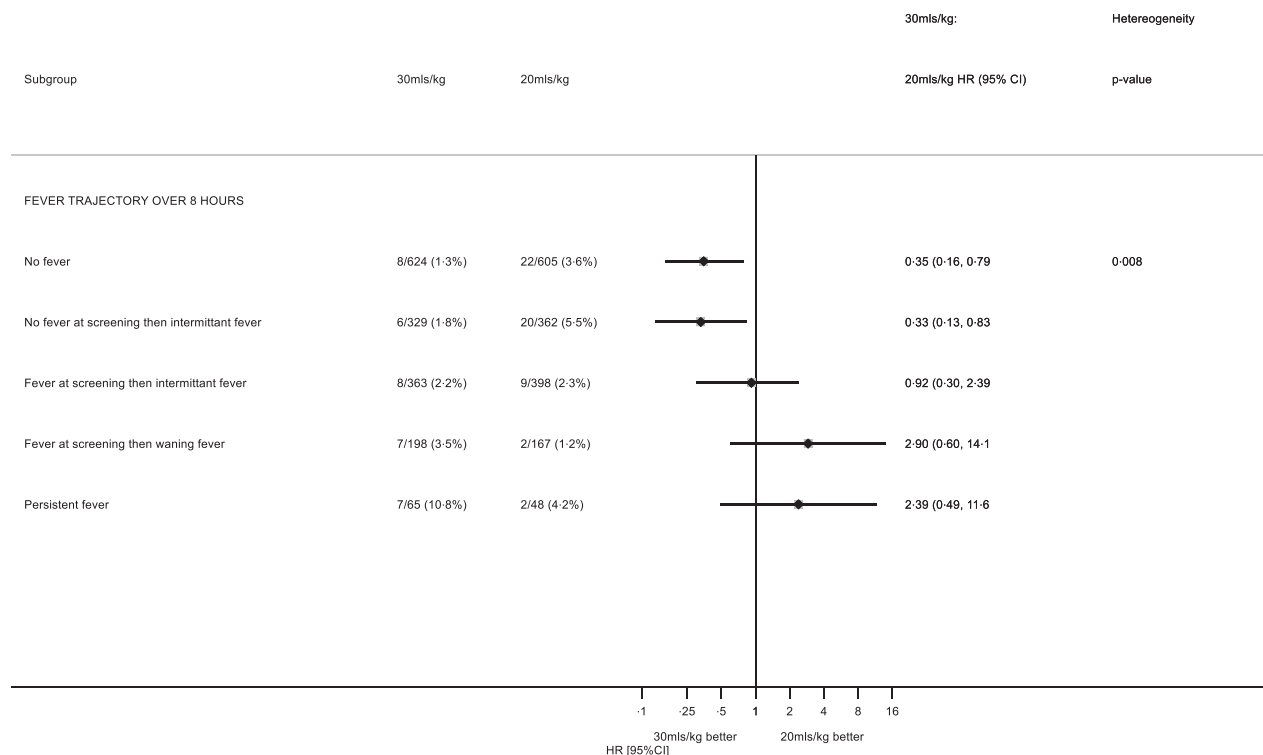


Fig 4. (A) Time to Transfusion in Immediate vs Control Comparison. (B) Cumulative incidence in second transfusion in the 30 mls/kg vs. 20 mls/kg split by fever and no fever. Note: A competing risks analysis with discharge, absconding and death as competing events. There are 39 transfusions given after 120 h from baseline not included on the graph. [Colour figure can be viewed at wileyonlinelibrary.com]

Table II. What were the indications for second transfusions by volume randomisation and by fever status?

Indication (at any time during admission)	Second transfusions for those randomised to transfusion arms, <i>n</i> (%)				
	Overall (<i>n</i> = 496 [†])	20 ml with fever* (<i>n</i> = 123)	20 ml without fever (<i>n</i> = 177)	30 ml with fever* (<i>n</i> = 83)	30 ml without fever (<i>n</i> = 113)
New Hb <40 g/l	185 (37)	45 (37)	82 (46)	18 (22)	40 (35)
New severity signs [‡] and Hb 40–60 g/l	9 (2)	3 (2)	1 (<1)	2 (2)	3 (3)
Known SCD or haemoglobinuria and Hb 40–60 g/l	206 (42)	56 (46)	63 (36)	37 (45)	50 (44)
Not per protocol [§]	96 (19)	19 (15)	31 (18)	26 (31)	20 (18)

Hb, haemoglobin; SCD, sickle cell disease.

*Axillary temperature $\geq 37.5^{\circ}\text{C}$.[†]497 transfusions were reported in the original trial paper but following publication it was found that one of those transfusions was of a negligible amount.¹⁴[‡]Severity sign (impaired consciousness and or increased work of breathing).[§]No recorded severity signs (impaired consciousness or increased work of breathing), no SCD or haemoglobinuria and Hb ≥ 40 and < 60 g/l.NB: excludes children who died prior to 8 hours (*n*=36) and 1 child that absconded. Time at risk from 8 hours onwards.**Fig 5.** Justification for transfusion volume management based on initial temperature and not on subsequent temperatures.

they to receive the higher volume. Although more complex, the TRACT showed that this approach could substantially reduce the demand for paediatric transfusion because the higher volumes given to children without fever would also significantly reduce the number of additional transfusions required. Our previous cost-effectiveness analysis indicated substantial savings for health services, with 60% reduction in blood transfusion required compared to immediate transfusion for all children with uncomplicated severe anaemia (Hb

<60 g/l).¹³ The unadjusted cost-effectiveness analyses, presented in the trial reports, demonstrated that the 'no immediate transfusion strategy' would cost \$66.46 (American dollars) per child in comparison to \$72.09 per child for an immediate transfusion strategy: the Meeting attendants recognised that the cost savings would be even greater where per-unit costs of donor blood are higher.

It was recognised that implementation of the algorithm in a real-life setting is required, in order to establish feasibility,

impact on mortality and whether it reduces unwarranted requests for paediatric transfusions. However, most paediatricians felt that the proposed recommendations for Hb monitoring could be a substantial hurdle for clinical services in some resource-limited hospitals (see accompanying Perspective Paper). As demonstrated in the cost-effectiveness models, major costs to health services are the 'hotel costs' (days in hospital) and the price of a blood transfusions. Total costs could be reduced by access to 24-h point-of-care Hb measurements, which would require advocacy to the health services and policy makers from both paediatricians and blood transfusion services.

The strengths of the TRACT trial include its size, multi-centre nature (Uganda and Malawi), broad eligibility criteria that enhances its generalisability and enables incorporation of large subgroups with malaria and SCD. In children with uncomplicated severe anaemia, we found a large group of children with undiagnosed SCD. We found no evidence that not transfusing these children (as they did not develop severity features or profound anaemia) resulted in a worse outcome (mortality to Day 28 and Day 180). On this basis, children with either diagnosed or undiagnosed SCD without severity features can be safely managed using a triggered transfusion strategy and thus incorporated into the algorithm.

As noted by the meeting stakeholders most transfusions in the TRACT were given very shortly after screening/identification of a case of severe anaemia (when axillary temperature was recorded), which may not be replicable in usual clinical practice. The relevance of a temperature taken hours before a blood transfusion becomes available was explored by additional analyses. These supported the recommendation that at the time that blood is ordered for a child who has no fever at that time, even if they subsequently developed an intermittent fever (Fig 5), it was still safe (and beneficial) to transfuse with a higher volume. Similarly, for children who have a fever at the time blood is ordered and which then wanes, there is no evidence to support that the requested 20 ml/kg volume would lead to any difference in outcome compared to 30 ml/kg. For children with a persistent fever prescribing 20 ml/kg is also supported by these analyses.

Conclusions

The new evidence provided by the TRACT could lead to important refinements of the WHO transfusion guidelines. First, standardising the definition of severe anaemia, we suggest an Hb <60 g/l. Second, clarifying the definitions of uncomplicated severe anaemia, and severe and complicated anaemia, by using clinical signs of severity. Third, a strong recommendation (based on high quality of evidence) that children with uncomplicated severe anaemia do not require immediate transfusion, irrespective of the underlying causes, but do require monitoring because ~50% will develop severe and life-threatening anaemia requiring subsequent

transfusion. Fourth, new recommendations to include a minimal number of clinical and Hb monitoring reviews in a child with severe anaemia (both uncomplicated and complicated) in order to identify those needing an initial or additional transfusion. Fifth, new recommendations on the volume of blood to transfuse, according to fever status when a blood transfusion is ordered, and that either WB or packed RCC (at half the volume) can be used. Finally, the guidelines should make it clear that there is no need for separate recommendations for children with malaria (even those with high parasite burdens), SCD or poor nutritional status. As the trial demonstrated, a high proportion of children presenting with severe anaemia in many of our present study sites had undiagnosed SCD. The group recommended, therefore, that admission to hospital with severe anaemia should prompt clinicians to test for SCD so that long-term follow-up and infection prophylaxis can be put in place.

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Patient and Public Involvement

Patients and public were not involved in the meeting or the interpretation of the results and development of the transfusion algorithm. We thank all the participants and their families, and staff from all the centres participating in the TRACT trial.

Supporting Information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Fig S1. Transfusion volume 30 *versus* 20 ml/kg comparison by fever status at screening and sickle cell disease status.

Fig S2. Stability of temperature following screening/randomisation.

Fig S3. Heart rate and respiratory rate over time from beginning of first transfusion.

Table SI. Suspected overload events: endpoint review.

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Appendix 1

Full list of Stakeholder Meeting Attendees

Name	Organisation	Role	Country
Dr Deogratias Munube	Mulago Hospital	President UPA	Uganda
Dr Bodo Bongomin	WHO Uganda	WHO Technical Officer	Uganda
Dr Eva Nabawanuka	Mulago National Referral Hospital	Study co-ordinator	Uganda
Dr Robert Opoka	Mulago Hospital	Co-PI TRACT trial	Uganda
Prof Peter Olupot Olupot	Mbale Regional Referral Hospital	Co-PI TRACT trial	Uganda
Dr Florence Aloroker	Soroti RRH	Co-PI TRACT trial	Uganda
Prof Philippa Musoke	MUJHU	TRACT DSMB	Uganda
Dr Ritah Nasiima	St Francis Hospital Nsambya	Treasurer UPA	Uganda
Prof Kathryn Maitland	KEMRI Wellcome Trust Research Programme	PI TRACT trial	Kenya
Hellen Mnjalla	KWTRP	TRACT trial manager	Kenya
Christabel Mogaka	KWTRP	Data Manager	Kenya
Prof Thomas Williams	KWTRP	TRACT co-investigator	Kenya
Dr Abubakarr Bah	Ola Daring Children's Hospital, Freetown	Head of Paediatrics Emergency Unit	Sierra Leone
Prof Elizabeth Molyneux	Liverpool School of Tropical Medicine	TSC Chair/ ETAT Lead	UK
Dr Christian Umuhoza	Secretary-General	Rwanda Paediatric Association	Rwanda
Dr William K. A. Obeng	Korle Bu Teaching Hospital Accra	ETAT Trainer	Ghana
Dr Charlyne Kilba	Korle BU Teaching Hospital, Accra	Critical care specialist	Ghana
Dr John Appiah	Paediatric Critical care	Senior specialist	Ghana
Dr Ismail Ticklay		ETAT Harare	Zimbabwe
Dr Yamikani Chimalizeni	UNIMA College of Medicine	TRACT trial Investigator	Malawi
Prof Russel Ware	Cincinnati Children's Hospital	Sickle cell disease Specialist	United States
Dr Roberta Petrucci	Medicines Sans Frontiere	International Paediatric Working Group	Geneva
Prof Diana Gibb	MRC Clinical Trials Unit at UCL	Professor of Epidemiology	UK
Prof Sarah Walker	MRC/CTU at UCL	Professor of Medical Statistics and Epidemiology	UK
Annabelle South	MRC/CTU at UCL	Policy and Research Impact Co-ordinator	UK
Prof Dora Mbanya	African Society for Blood Transfusion (AfSBT)	AfSBT President	Cameroon
Dr Sophie Uyoga	KWTRP	TRACT trial Investigator	Kenya
Dr Bridon Mbaya	Malawi Blood Transfusion Service (BTS)	Medical Director	Malawi
Dr ET Mberi	Zimbabwe National BTS	Director National Blood Transfusion Centre	Zimbabwe
Dr Dorothy Kyeyune	Uganda BTS	Director	Uganda
Dr Claude T. Tagny	Haematology and Transfusion Service, University of Yaounde	Director National Blood Transfusion Centre	Cameroon
Dr Saliou Diop	Senegal BTS	Director National Blood Transfusion Centre	Senegal
Dr Faten Moftah	AfSBT	AfSBT member	Egypt
Dr Michael E. Acquah	Ghana BTS	Director	Ghana
Prof Philip Olatunji	AfSBT	Professor of Haematology/Vice President, AfSBT	Nigeria
Dr Magdalena Lyimo	Tanzania BTS	Director	Tanzania
Prof Ludovic Anani	Benin BTS	Former manager, Blood Transfusion Agency of Benin	Benin
Dr Shirley O. Ofori	Kumasi BTS	Director	Ghana
Dr Charles Engoru	Soroti RRH	TRACT trial investigator	Uganda
Prof Imelda Bates	Liverpool School of Tropical Medicine	TRACT trial investigator	UK/TRACT trial

AfSBT, Africa Society for Blood Transfusion; BTS, blood transfusion services; CTU, Clinical Trials Unit; DSMB, Data and Safety Monitoring Board; ETAT, Emergency Triage Assessment and Treatment; KWTRP, Kenya Medical Research Institute Wellcome Trust Research Programme; MRC, UK Medical Research Council; MSF, Médecins Sans Frontières; MUJHU, The Makerere University - Johns Hopkins University Research Collaboration; PI, principal investigator; RRH, Regional Referral Hospital; STM, School of Tropical Medicine; TRACT, Transfusion and Treatment of severe anaemia in African Children Trial; TSC, Trial Steering Committee; UNIMA, University of Malawi; UPA, Uganda Paediatric Association; WHO, World Health Organization.