A Bleeding Nightmare - Improving Patient Outcomes Paediatric Major Haemorrhage



Dr F Chowdhury Consultant Haematologist in Transfusion Medicine Honorary Senior Clinical Lecturer- Imperial ICHNT / NHSBT Post Graduate Education Centre, East Surrey Hospital 8th March 2019

The World of Paediatrics







HOME GUIDELINES GUIDELINES TRANSFUSION FOR FETUSES, NEONATES AND OLDER CHILDREN

Transfusion for Fetuses, Neonates and Older Children

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Appropriate transfusion of fetal and paediatric patients of all ages is vital in order to balance transfusion benefits against risks. These risks include transfusion of an incorrect blood component due to errors such as mistaken patient identity, or unpredictable acute transfusion reactions (Stainsby et al, 2008). Recent studies suggest that a significant percentage of paediatric transfusion recipients receive only one transfusion during their admission (Slonim et al, 2008; New et al, 2014), raising the possibility that some may be avoidable. Specialised components are available for transfusion to different paediatric patient groups and for different clinical indications.

Plasma components have been imported for all patients born on or after 1st Jan 1996 in order to reduce the risk of transfusion transmission of variant Creutzfeldt-Jakob disease (vCJD; see section 7). Additional component safety measures are applied for fetal and neonatal patients, who are particularly vulnerable recipients because of their small size and developmental immaturity and who also have the longest potential lifespan. The clinical section focuses largely on aspects relating to transfusion indications and administration, whereas the laboratory section contains most of the information relating to pre-transfusion testing and component selection.

Definitions

Neonate <28 days old

Infant 28 days – 1 yr

Child 1-18 yr (but >16 admitted under adult ward)

PAEDIATRIC MAJOR HAEMORRHAGE PROTOCOL Rapid blood loss with shock or with no likelihood of control. Anticipated or actual administration of 40 mL/Kg of blood FOR USE IN CHILDREN under 50Kg Larger children - use adult protocol

Call 2222. State "Major Haemorrhage". Give Hospital and Location

Nominated Transfusion Co-ordinator MUST CONTACT Blood Transfusion with the following:

- 1. Patient identification Hospital Number, Name & Date of Birth (or trauma patient details)
- 2. Approximate weight of child
- 3. Patient location
- 4. Name and contact details of Nominated Transfusion Co-ordinator for on-going communication
- 5. Cause of bleeding
- 6. Confirm Group &Screen, Full Blood Count & Coagulation Screen samples are being sent to laboratories - If difficulty obtaining blood samples ensure 4mL EDTA sample sent for crossmatch as a priority

Call the Blood Transfusion Laboratory Monday-Friday 9am-5pm Ext. 21157 Trauma Ext. 22043 Bleep 1611

Definition of Massive Blood Loss 80mls/kg in 24 hours 40mls/kg in 3 hours 2-3mls/kg/min

Platelets

Immediate if on site

The Blood Transfusion Laboratory will issue: Availability of Blood For Collection 20ml/kg O negative RBC & 20ml/kg FFP Or 20ml/kg group specific* RBCs & 20ml/kg FFP (*if valid sample in Laboratory) Fresh Frozen Plasma (If no valid samples continue to issue emergency blood) 30 minutes to thaw Once these components are collected from the laboratory: The laboratory will continue to issue 20ml/kg RBCs and 20ml/kg FFP. whilst the patient is bleeding, until stood down from MHP Cryoprecipitate Administer Tranexamic Acid 30 minutes to thaw At this stage consider requesting: Platelets 15-20ml/kg (up to 1 pack)

- Cryoprecipitate 10ml/kg (up to 10 single-donor units)
- After 10 RBCs given consider Fibrinogen concentrate (50mg/kg)

Porters:

- Report to Transfusion Lab to collect blood, then to ward, except for:
- A&E at SMH & CXH: Report to A&E whereby staff will tell porters when to collect blood components

The clinical area will:

- Nominate a Transfusion Co-ordinator to ensure effective management of blood components
- Send full blood count & coagulation screen samples as a baseline and hourly thereafter
- Send repeat group & save sample if requested
- Ensure the patient's Consultant has been informed (if not already aware)
- Discuss on-going management including authorisation of other clotting factors with the Haematology SpR
 (contact through Switchboard if contact details not known)
- Where required ensure RBC are warmed appropriately
- Inform the Blood Transfusion Laboratory when STOOD DOWN

Imperial College Healthcare NHS Trust PAEDIATRIC MAJOR HAEMORRHAGE PROTOCOL 2018 Version 7 Please remove and destroy any previous versions of the Paediatric Major Haemorrhage Proto

• Call 222

- State hospital / location
- Call transfusion lab
- Provide patient details
- Maintain contact with lab throughout MH call
- GIVE TXA EARLY

Remember to STAND DOWN

What if peripheral access is not possible?







What is the total blood volume for a 2 year old?

Define massive blood loss

How would you manage this case and what product support would you recommend?

4. What are your targets?

1. Total volume 80mls/kg

2. Massive blood loss

- 80mls/kg in 24 hours
- 40mls/kg in 3 hours
- 2-3mls/kg/min



Activate Resuscitation



Immediate transfusion:

Early use of FFP, CRYO and platelets prior to the results of coagulation tests where bleeding is on-going.

After initial resuscitation, appropriate aliquots to be transfused are as follows:

> RBCs 20 mls/kg aliquots (O D negative or ABO & D specific)

FFP in 20 ml/kg aliquots 10ml/kg

Cryoprecipitate

Platelets in 15-20mls aliquots – consider after every 40 ml/kg RBCs

20 ml/kg RBCs O D-negative or ABO Rh D compatible

1 FFP:2 RBC

A case of an unusual complication

Transfusion Associated Hyperkalaemia

Recognised complication of massive RBC in Paediatrics. 8/9 case reports (in English) of TAHCA. K+ ranged from 9.3-12mmol/L. Rate of blood transfusion, more so than the total volume, cardiac output and site of infusion were key in development of transfusion-associated hyperkalaemia in paediatric patients. (Lee *etal* (2014)

Often multifactorial & difficult to isolate completely to transfusion

RBC during storage from 0-42 days studied by Sezdi, Bayik, Ulgen (2006)

Vraets *etal* (2011) cited that the increase in potassium appears to be roughly linear with time.

Weiskopf *etal* (2005) studied the effects of irradiation (25 Gy) and washing on potassium:

Potassium of irradiated units increaese more rapidly than non-irradiated units. Potassium of units washed after irradiation increased more slowly than if washed before irradiation.

Vraets etal 2011

Irradiated and then washed RBC

0 hours	1.6 +/- 0.3mmol/L
6 hours	5.3 +/- 0.5mmol/L
12 hours	8.6 +/- 1.0 mmol/L
24 hours	14.3 +/- 1.3mmol/L
Non irradiated RBC (washed)	
24 hours	5.9 +/- 1.4 mmol/L

Causes of high K in this case



? Cold

units

Hypovolaemic shock

Storage age

Lack of washing

?Rapid rate of transfusion

Measures to reduce K+

Anticipate and replace blood loss before significant haemodynamic compromise

Use large bore (>23gauge) peripheral access

Check and correct electrolyte abnormalities frequently

Use fresher RBC in massive transfusin

Changes in Practice Locally

Changes to practice already implemented in the transfusion lab

1. Avoid issuing irradiated RBC units in all major haemorrhage / trauma calls

 Ensure that mixture of RBC units issued: ratio of roughly 2 units >21 days with 2 units <14 days old.

> 3. Reduced stock holding of irradiated RBC units for BMT cover in laboratory at any one time to minimise risk of mixing irradiated units with general stock.

Changes recommended in clinical setting for implementation

 Clearly document time each unit given to facilitate look backs using form Blood component administration records for Massive Haemorrhage (see attached document)

> 2. Infuse blood through cell salvage which has the effect of washing the RBC thereby reducing the potassium load given to the patient.

References

Lee A, Reduque L, Luban N, Ness P, Anton B Heitmiller E (2014) Transfusion-associated hyerkalaemic cardiac arrest in paediatric patients receiving massive transfusion. <u>Transfusion</u>. 2014 Jan; 54(1):244-54Tranfusion

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