

The Development of a Regional Guideline for Management of Massive Haemorrhage

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Background

- RTC members requested Regional Guideline to address new developments in this field:
 - Experience from the military arena on use of ‘haemostatic resuscitation’^{1,2}
 - Near patient testing techniques such as thromboelastography³
 - Use of additional therapies such as tranexamic acid⁴, prothrombin complex concentrate⁵ and fibrinogen concentrate⁶



Background

- The National Patient Safety Agency has raised concerns with regards to delays in availability of blood in emergencies. Over a three year period (October 2006-Oct 2009) 10 deaths and 46 incidents associated with major morbidity were reported
- Emphasis needs to be on rapid provision of blood and blood components
- A key element is the effective communication between all staff who will be involved in the provision and transportation of blood



A toolkit or a guideline?

- There is a paucity of good randomised controlled trial evidence on which to base guidelines – most publications describe level III or IV studies
- Two groups actively producing up to date guidelines: the Association of Anaesthetists of Great Britain and Ireland (AAGBI) and Australian Patient Blood Management Program – both currently completing a consultation process on a final draft
- Decision to develop a toolkit rather than a guideline as a means of putting current evidence into practice



Components of the Toolkit

- A simple algorithm for the management of major haemorrhage based on current evidence that can be customised according to local circumstances (Trust or Specialty Specific)
- Specialties covered:
 - Major trauma General and vascular surgery
 - Cardiac surgery Obstetrics
 - Paediatrics Gastrointestinal haemorrhage
- A section on effective communication
- A section on monitoring the management of major haemorrhage



Who is the toolkit for?

- For hospital transfusion teams and committees.
- It is recommended that the algorithm is used as a template for the production of a flow chart in each Trust that will complement the Trust guideline(s) for the management of major haemorrhage



Who is the management algorithm for?

- The junior doctor / senior nurse who may be the first person to see the patient and must be able to recognise the early stages of a major haemorrhage situation and know when and who to call for support
- The senior staff called as part of the emergency response team
- The laboratory and supporting services (eg: portering)



Method

- Call for volunteers via RTC membership Nov 2009
- Literature review performed Jan 2010
- Volunteers divided into 8 subgroups
- Lead from each subgroup on steering group
- First steering group meeting held 23/4/10



The subgroups

- **Communications and logistics**
- **Near patient testing, haemostasis and lab issues**
- **Obstetric haemorrhage**
- **Major Trauma**
- **Gastro intestinal haemorrhage**
- **General / vascular surgery**
- **Paediatrics**
- **Cardiac surgery**



The process

- Interim workshop May 14th
- Final workshop 18th June
- Final draft circulated to RTC members, critical care, trauma and vascular surgery networks in NW consultation July – September 2010



Dilemmas

- Is it possible to fit everything on one page in a clear and simple format?
- Is pathway just for transfusion management or should it include advice on resuscitation and other measures to control haemorrhage?
- Definition of major haemorrhage and when to trigger pathway
- A separate pathway for each specialty or one pathway for all?
- Major haemorrhage packs ('shock packs') or not? Just for major trauma, or applicable to all major haemorrhage situations?
- Thawed AB plasma on standby or group specific plasma when group known?
- When to order platelets and how many doses?
- The role of TEG / near patient testing
- The role of tranexamic acid, fibrinogen concentrate, prothrombin complex concentrate (other than for reversal of warfarin), use of rFVIIa in light of SPC advice against using the product in this situation



Pros and Cons of Formula driven Massive Transfusion Protocols

Pros

- Reduce mortality from bleeding
- Improve speed of delivery of blood components
- Decrease need for communications back and forth between clinical area and lab
- Prevent onset of coagulopathy
- Reduce dependency on lab testing in acute resuscitation phase

Cons

- Based on level III and IV evidence mainly in major trauma
- Exposure to additional units of FFP and platelets will increase risk of complications such as TRALI, organ failure, thrombosis and sepsis
- Inappropriate triggering of use of formula driven care in non massive transfusion patients
- Increased wastage of FFP and platelets
- Depletion of platelet and plasma stocks



Draft matrix

Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
<p>Trigger for activation Patient bleeding / collapses Ongoing severe bleeding (overt / covert) eg: 150 mls / min and not controlled With Class III / IV shock and / or coagulopathy Or if bleeding ongoing after 4 units of blood and patient is haemodynamically unstable due to blood loss</p> <p>Paediatric version Ongoing severe bleeding (overt / covert) and received 20ml/kg of red cells or 40ml/kg of any fluid for resus in preceding hour With hypovolaemic shock and / or coagulopathy</p>	<p>For population by subgroups: Specific points re clinical situations that help to identify major haemorrhage early</p>	<p>Resuscitation Assessment High flow oxygen Large bore IV access Invasive BP and CVP monitoring if possible Maintain normothermia and avoid acidaemia Warm IV fluids, use blood warmer and level 1 infuser ‘Hypotensive resuscitation’ until haemostasis achieved (in absence of head injury)</p> <p>Stop the bleeding Simple measures: Direct pressure / tourniquet if appropriate Surgery Interventional radiology Endoscopic techniques</p>	<p>Who is responsible for activation? Depends on the clinical situation – for subgroup suggestions Toolkit should be designed to allow junior doctor, senior doctor or senior nurse / midwife to recognise major blood loss situation and activate pathway (incorporate use of EWS)</p>




Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
Activation of pathway Assemble emergency response team (consultant delivered care recommended) <ul style="list-style-type: none"> ▪ Team leader ▪ Team roles: <ul style="list-style-type: none"> ▪ ABC (Airway, breathing, circulation) ▪ Sample taker / investigation organiser / documenter ▪ Communicator ▪ Transporter (may be member of clinical team or porter) ▪ Theatre team / other team members as required 	<p>How team is mobilised and who is in team may vary between clinical groups</p> <p>Each subgroup to consider who should be notified and how</p> <p>Remember to include: laboratory person, portering team, on call consultant haematologist / SpR, consider cell salvage operator (esp if obstetric, elective surgery, vascular emergency, major trauma in theatre)</p>	<p>Ongoing assessment and resuscitation</p>	<p>Contact switch to coordinate emergency response team:</p> <p>'Major Haemorrhage'</p> <p>'Location'</p> <p>Specialty</p> <p>Alert: emergency response team</p>



Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
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<p>Decision on immediate blood requirements</p> <p>Team leader to decide on immediate blood requirements :</p> <ul style="list-style-type: none"> •immediate use O Neg •30 – 60 mins Group Specific •60-90 mins Crossmatched 	<p>Pathway should be populated locally with sites of O Neg storage and number of units available and realistic time for blood availability for group specific and crossmatched blood, depending on transit arrangements and lab time</p>	<p>Resuscitation: permissive hypotension, oxygenation, normothermia, correction of acidosis etc</p> <p>Turning off the tap: surgery, radiological / endoscopic control</p> <p>Cell salvage</p>	<p>Communicate with lab ext : (out of hours:)</p> <p>Communication should only occur between the designated communicator and the BMS</p> <p>•‘This relates to major haemorrhage situation’</p> <ul style="list-style-type: none"> •Name, location and ext no. •The patient’s details: surname, first name, Patient ID number, DOB or gender / A+E number if unidentified •Whether O Neg has been used and how much •Order MHP Pack 1 ASAP – see below •Request urgent investigations – see below
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Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
Blood sampling and other tests U+E, FBC, Crossmatch, PT, APTT, Fibrinogen, ABG, Calcium, lactate Other investigations	Subgroups – please populate with tests that are required	Resuscitation, permissive hypotension, oxygenation, normothermia, correction of acidosis etc Turning off the tap: surgery, radiological / endoscopic control	Communication to lab that urgent samples are on their way – see above – need appropriate transport Lab needs to phone results back to clinical area (to ‘communication lead’)
Near patient testing TEG / ROTEM / near patient PT	Subgroups – please consider role in your specialty and hospital variation		
Choice of components Massive Haemorrhage pack one (MHP 1) 4 units red cells 4 FFP 1 dose platelets (lab may order 2 doses of platelets on blue light)	Subgroups – are there any variations to MHP 1 Also consider variation between hospitals – if platelets in stock then may not need to order 2 packs up front	Use of calcium after 5 units red cells – specify drug dose and method of admin Tranexamic acid (major trauma 1g over 10 mins followed by infusion of 1g over 8 hours), also has role in cardiac and vascular surgery Cell salvage	Contact with lab to order MHP 1 (in reality this is likely to be done earlier in pathway – see decision on immediate blood requirements) Lab orders two doses of platelets and rings communication lead to let clinical team know when blood components are ready
Paediatric version Order MHP 1 Administer up to: Red cells 40 ml/kg FFP 20 ml/kg Platelets 10 ml/kg			

Paediatric Major Haemorrhage pack 1 (MHP 1) order these volumes, volumes for administration are detailed in the flow chart

Weight	Red cells	FFP	Platelets
< 5kg	2 Paediatric Units (80-100ml)	2 'neonatal' units of Methylene Blue (MB) treated FFP (100ml)	1 paediatric pack of platelets (50ml)
5 -10kg	1 Adult unit (250ml)	1 paediatric unit MB treated FFP (225ml)	2 paediatric pack of platelets (100ml)
10 – 20 kg	2 Adult units (500ml)	2 paediatric units MB treated FFP (450ml)	1 Adult apheresis pack (200ml)
20 – 40 kg	4 Adult units (1000ml)	4 paediatric units MB treated FFP (900ml)	1 Adult apheresis pack (200ml)



Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
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Repeat blood tests PT, APTT, fibrinogen, FBC, U+E, Ca, lactate Near patient tests may be more timely in guiding therapy	Subgroups – please populate with tests that are required	Cell salvage if available and appropriate Consider ratios of other products 1 unit of red cells = approximately 250 mls salvaged blood	Communication to lab that urgent samples are on their way – see above – need appropriate transport Lab needs to phone results back to clinical area
Target results to aim for in ongoing bleeding situation: Suggested figures: Hb 10 g/dl Platelets $75 \times 10^9/l$ Fibrinogen $> 1 \text{ g/l}$ PT ratio < 1.5 APTT ratio < 1.5	Subgroup to refine suggested figures depending on clinical scenario: Eg: ?fibrinogen $> 2\text{g/l}$ in obstetric haemorrhage; platelets $> 100 \times 10^9/l$ in head injury		



Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
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<p>If bleeding ongoing: Request MHP 2 4 red cells 4 FFP 1 dose platelets (will have already been ordered with MHP 1) 2 packs cryoprecipitate</p>	<p>Subgroups – are there any variations to MHP 2? Also consider variation between hospitals</p>		<p>Contact lab to order MHP 2</p>
<p>Paediatric version Order MHP 2 Administer up to: Red cells 20 ml/kg FFP 20 ml/kg Platelets 10 ml/kg Cryoprecipitate 10 ml/kg</p>			
<p>Repeat blood tests PT, APTT, fibrinogen, FBC, U+E, Ca, lactate Near patient tests may be more timely in guiding therapy</p>	<p>Subgroups – please populate with tests that are required</p>		<p>Communication to lab that urgent samples are on their way – see above – need appropriate transport Lab to phone results</p>




Paediatric Major Haemorrhage pack 2 (MHP 2) order these volumes, volumes for administration are detailed in the flow chart

Weight	Red cells	FFP	Cryoprecipitate	Platelets
< 5 kg	2 Paediatric Units (80-100ml)	2 'neonatal' units of Methylene Blue (MB) treated FFP (100ml)	1 single donor unit Mb treated (40ml)	1 paediatric pack of platelets (50ml)
5 – 10 kg	1 Adult unit (250ml)	1 paediatric unit MB treated FFP (225ml)	2 single donor units (80ml)	2 paediatric pack of platelets (100ml)
10 – 20 kg	2 Adult units (500ml)	2 paediatric units MB treated FFP (450ml)	1 pool (5 units) (200ml)	1 Adult apheresis pack (200ml)
20 – 40 kg	4 Adult units (1000ml)	4 paediatric units MB treated FFP (900ml)	2 pools (10 units) (400ml)	1 Adult apheresis pack (200ml)



Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
<p>If still bleeding: Contact haematologist Consider PCC, fibrinogen concentrate**, rFVIIa* Further red cells Tailor blood product use according to coagulation results *rFVIIa : use with caution according to local protocol where conventional measures to control bleeding have failed **PCC and fibrinogen: further recommendations to be made as evidence becomes available</p>	<p>Subgroups – please populate with specific requirements for your speciality</p>		<p>Communicate with haematologist</p> <p>Haematologist to review results and make recommendations – via communication lead on team</p>
<p>Additional haemostatic challenges: Warfarin: Check urgent INR, reverse anticoagulation with vitamin K 5-10mg IV and PCC according to local policy Aspirin / clopidogrel: consider additional dose of platelets in each MHP, discuss alternative approaches with haematologist Unfractionated heparin / LMWH / coagulopathy: discuss with haematologist</p>		<p>Storage and handling of blood components</p> <ul style="list-style-type: none"> •Red cell transfusion should commence as soon as practicable after removal from cold storage •Platelets must be kept at room temperature and given through a fresh giving set •FFP and cryoprecipitate must be infused ASAP •Please follow NW Transfer of Blood Policy if blood is to be / has been transferred with the patient to / from another Trust •Staff must be familiar with local protocols to record fate of units 	

Stage of pathway	Sub group variation or hospital variation	Other management aspects	Communication
Stand down Bleeding controlled Mechanism for checking / replenishing stocks of O Neg in satellite fridges Replenish laboratory stocks of red cells / FFP / cryo / platelets if appropriate	Subgroups: specify definition of controlled situation	Complete documentation Record of / Timings of blood samples and communications, transfusion documentation, traceability documentation, recording of vital signs Venous thromboprophylaxis should be commenced ASAP after bleeding controlled	Communicate with lab and transport staff etc that situation under control
Audit Monitor appropriateness of pathway activation Monitor blood supply, usage and wastage Monitor patient outcome data Review process and team structure Develop test exercises, drills Regular testing of initiation system	Consider how to test the pathway in different clinical scenarios		Whose role is this? – transfusion lab sends team leader audit proforma on stand down ?HTT to coordinate? Supervised and reviewed by RTC An audit proforma has been designed as part of the RTC toolkit
Incident reporting All incidents where there are delays or problems in the provision of blood in an emergency must be reported and investigated locally, and reported to the NPSA and the Serious Hazards of Transfusion (SHOT) scheme.	Consider who should create incident report Describe method in local trust with recommendation of level of incident and timescale for incident management		Communication of results of investigation to change practice – oversight of all incidents in HTC 

NHSBT logistics

1. Recognise trigger and activate pathway for management of massive blood loss; assemble the emergency response team

[REDACTED]

2. Allocate team roles

- I. Team leader
- II. Communicator – dedicated person for communication with other teams, especially the transfusion laboratory
- III. Sample taker / investigation organiser / documenter
- IV. Transporter - porter, member of team from clinical area) ,

[REDACTED]

3. Complete request forms / take blood samples, label samples correctly / recheck labelling

U+E, FBC, Crossmatch, PT, APTT, Fibrinogen, ABG, Calcium, lactate

[REDACTED]

4. Request blood / blood components – team leader should decide on use of:

- I. Emergency O Neg (immediate)

[REDACTED]

- II. Group specific **insert time to availability here**

- III. Crossmatch **insert time to availability here**

- IV. Communication with lab: contact BMS

[REDACTED]

and inform them:

- a. Your name, location and ext number
- b. 'this relates to the major haemorrhage situation'
- c. The patient's details ideally surname, forename, hospital number, DOB (insert acceptable details for unknown casualty here)
- d. Whether O Neg has been used and how many units
- e. Order major haemorrhage pack 1
- f. Contact lab if blood has been transferred in with patient from another trust

5. Transport samples to laboratory, BMS to ring back with results of urgent investigations and when blood / components are ready, collect blood and blood components from laboratory

[REDACTED] ated

6. Communicate stand down of pathway and let lab know which products have been used

7. Ensure documentation is complete

- I. Clinical area: monitoring of vital signs, timings of blood samples and communications, transfusion documentation in patient casenote record, return traceability information to laboratory
- II. Laboratory: keep record of communications / telephone requests in patient laboratory record



North West Regional Transfusion Committee
incorporating North Wales

Benefits of the process

- Good engagement between different specialities and disciplines
- Sharing of best practice from Trusts around the region
- Sense of ownership of the final document
- Good links between clinical users and NHSBT
- Group is small enough to reconvene to allow for regular updating of the document
- RTC will support regional audit



References



North West Regional Transfusion Committee
incorporating North Wales

Subgroup membership

Name	Position	Trust	Subgroup
Mike Desmond	Cardiac Anaesthetist	Liverpool Heart and Chest	cardiac surgery
Richard Williams	Cardiothoracic Surgeon	Liverpool Heart and Chest	cardiac surgery
Niall O'Keeffe	Cardiac Anaesthetist	Central Manchester	cardiac surgery
Kate Pendry	Consultant Haematologist	NHSBT / CMFT	communication and logistics
Emma Milser	Transfusion Practitioner	SMUH	communication and logistics
Tony Davies	Transfusion Liaison Practitioner	NHSBT	communication and logistics
John Tappin	Consultant Haematologist	St Helens	communication and logistics
Simon Cunningham	Consultant Obstetrician	Leighton	communication and logistics
Eithne Hughes	Transfusion Practitioner	Glan Clwyd	communication and logistics
Christine McQuillan	Transfusion Lab Manager	Aintree	communication and logistics
Tracey Scholes	Hospital Liaison Manager	NHSBT	communication and logistics
Gurvinder Banait	Consultant Gastroenterologist	WWL	gastrointestinal haemorrhage
Prasad Neeraj	Consultant Gastroenterologist	WWL	gastrointestinal haemorrhage
David Raw	Consultant Anaesthetist	Aintree	gen / vascular surgery
Dr Nagaraja	consultant anaesthetist	Aintree	gen / vascular surgery
Oliver Hill	Consultant Anaesthetist - Obstetric and Emergency	UHSM	gen / vascular surgery
Tushar Mahambrey	Consultant Anaesthetist / ICU	St Helens	gen / vascular surgery
Jane Uttley	Transfusion Lab Manager	Stockport	gen / vascular surgery
Winston de Mello	Consultant anaesthetist	UHSM	major trauma
Patrick Nee	Consultant A+E	St Helens	major trauma
Paul Wallman	Consultant A+E	Trafford	major trauma
Derek Pegg	Trauma and Orthopaedics	Leighton	major trauma
Andy Curran	Med Director	NW Ambulance	major trauma
Lilian Parry	Transfusion Lab Manager	St Helens	major trauma
Jackie McLennan		Wythenshawe	major trauma
Clare Barnes	Consultant Haematologist	Bolton	near patient testing, haemostasis and laboratory
Emma Searle	SpR Haematology	NHSBT	near patient testing, haemostasis and laboratory
Sharran Grey	TLM/TP	Bolton	near patient testing, haemostasis and laboratory
Lynne Mannion	Transfusion Practitioner	Blackburn	near patient testing, haemostasis and laboratory
Sarah Haynes	Autologous Transfusion Coordinator	SMUH	NPT / or gen / vascular surgery / obs
Dr Jothilakshmi	Consultant Obstetrician	Pennine Acute	obstetric haemorrhage
David Burch	Consultant Obstetrician	Lancaster	obstetric haemorrhage
Veera Gudimetla	Consultant Obstetrician	Leighton	obstetric haemorrhage
Shanthi Pinto	Consultant Obstetrician	Leighton	obstetric haemorrhage
Teresa Kelly	Consultant Obstetrician	Salford Royal	obstetric haemorrhage
Shuba Mallaiah	Obstetric Anaesthetist	Liverpool Women's	obstetric haemorrhage
Stephen Gilligan	Consultant Anaesthetist / ICU	Blackburn	or gen / vascular surgery
Bimal Mehta	Consultant Paed Emergency Medicine	Alder Hey	paediatric
Richard Craig	Consultant Paediatric Anaesthetist	Alder Hey	paediatric
Denise Bonney	Paediatric Haematologist	RMCH	paediatric
Wendy Ogden	Transfusion Lab Manager	RMCH	paediatric
Srivedi Kuchi	Consultant Paediatric Anaesthetist	Alder Hey Children's Hospital	paediatric
Sonia Desilva		St Helens	burns surgery



Steering Group membership

Group	Name of lead
Communications and logisitics	Tony Davies
Near patient testing, haemostasis and lab issues	Clare Barnes
Obstetric haemorrhage	Shuba Malliah
Major Trauma	Winston de Mello
Major trauma	Tushar Mahambrey
Gastro intestinal haemorrhage	Gurvinder Banait
Paediatrics	Bimal Mehta
Paediatrics	Denise Bonney
Cardiac surgery	Mike Desmond
Gen vascular surgery	Dr Nagaraja
Gen vascular surgery	David Raw
A+E	Paul Wallman
Burns surgery	Sonia Desilva
Other members:	Sarah Haynes
	Kate Pendry

