

Trust Standard Operating Procedures (SOP)

Management of Massive Haemorrhage

Document Control

Author/Contact	Consultant Haematologist Dr David Simcox	
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Management of Massive Haemorrhage

Appendix 4 of the EQMS 5 V10 Blood Transfusion Policy

Steps for Successful Coordination in Massive Haemorrhage

4.1 Initial Action

Recognise that patient has a massive haemorrhage and Senior Medical staff activate pathway for management of massive haemorrhage; Communication lead to activate Massive Haemorrhage pathway by calling “2222” and stating “Massive Haemorrhage” followed by the location and contact extension number. The laboratory will contact the clinical area back on this number.

4.2 Allocate team roles (Clinical Area)

- Team leader
- Communication lead– dedicated person for communication with other teams, especially the Transfusion Laboratory and support services. Should this change during the event the laboratory MUST be inform as soon as possible.
- Sample taker / investigation organiser / documenter
- Transporter - porter, member of team from clinical area

4.3 Complete request forms / take blood samples, label samples correctly / recheck labelling U+E, FBC, Crossmatch, PT, APTT, Fibrinogen, ABG, Calcium, Lactate

All blood request forms must be completed as per Trust policy - Laboratory Investigations Minimum Data Standard.

4.4 Communication lead to contact laboratory:

Massive Haemorrhage Activation - 2222

Transfusion (RLUH) Ext 4331 / 4332 24 hours

Inform the Switchboard Operator:

- Massive Haemorrhage
- State Location
- State Contact Extension Number

The Biomedical Scientist (BMS) will call back on this extension number and will require:

- Your name, location and extension number
- The patient’s details: ideally surname, forename, hospital number, DOB (For unknown patients, as a minimum they must have a unique hospital unit number. For any unknown details write ‘unknown’ do not leave blank spaces).
- Order massive haemorrhage pack(s) and /or appropriate blood components (See step 4.5 for guidance on timings)
- Contact the Transfusion Laboratory if blood has been transferred in with a patient from another Trust or a patient is being transferred to another Trust.

BMS in Transfusion Laboratory will complete part 1 of the Massive Haemorrhage Form (Appendix 4) using the information given above and ensure part 2 is made available to be taken to clinical area.

4.5 Request blood / blood components

Team leader should consider **Massive Haemorrhage Pathway (Appendix 1)** and decide on use of:

- **Red Cells**
- **Emergency Group O*** (immediate) Available within 5 minutes of call to Laboratory.
**O Negative to females of child bearing potential. O Positive to females not of child bearing potential and all males.*
- **Crossmatched**
Available usually within 45 minutes of sample arriving in Laboratory providing the antibody screen is negative (90% of cases). The Transfusion Laboratory will inform the clinical lead if there will be any delay due to a positive antibody screen, anomalous grouping or technical problems.
- **FFP and Cryoprecipitate:**
Be aware that FFP and Cryoprecipitate will take approx. 30 minutes from request to thaw and be issued. If patients' blood group is unknown group AB FFP/Cryoprecipitate will be issued. Once the patients group is known group specific frozen components will be issued.
- **Platelets**
It should be noted that it **cannot be guaranteed** that the Transfusion Laboratory will have a stock of platelets. This is governed by the availability of national supplies. Platelets may need to be ordered from the NHSBT. **The minimum time for platelets to be delivered in an emergency situation is 65 minutes.**
For patients who group is unknown group A platelets will be issued if available. Once the patients group is known group specific platelets will be issued.
If there are NO platelets available within the Laboratory, for patients whose group is unknown, group A platelets will be ordered from the NHSBT and issued. Once the patients group is known group specific platelets will be ordered from the NHSBT issued.
The policy does not advocate the availability of thawed group AB FFP and group A platelets on standby.

Repeat process as clinical situation requires further blood components – Please refer to Massive Haemorrhage Pathway for guidance (Appendix 1) It should be understood that the Massive Haemorrhage Pathway is intended as guidance and that further blood components can be ordered by senior Anaesthetists, Intensivists and ED medics should the clinical situation warrant it.

4.6 The Clinical / Laboratory interface

- Communication lead to arrange for transport of samples / request form to the laboratory
- BMS to telephone communication lead with results of urgent investigations
- BMS to telephone communication lead when blood / blood components are available for collection.
- Communication lead to arrange to collect blood and blood components from the Laboratory.

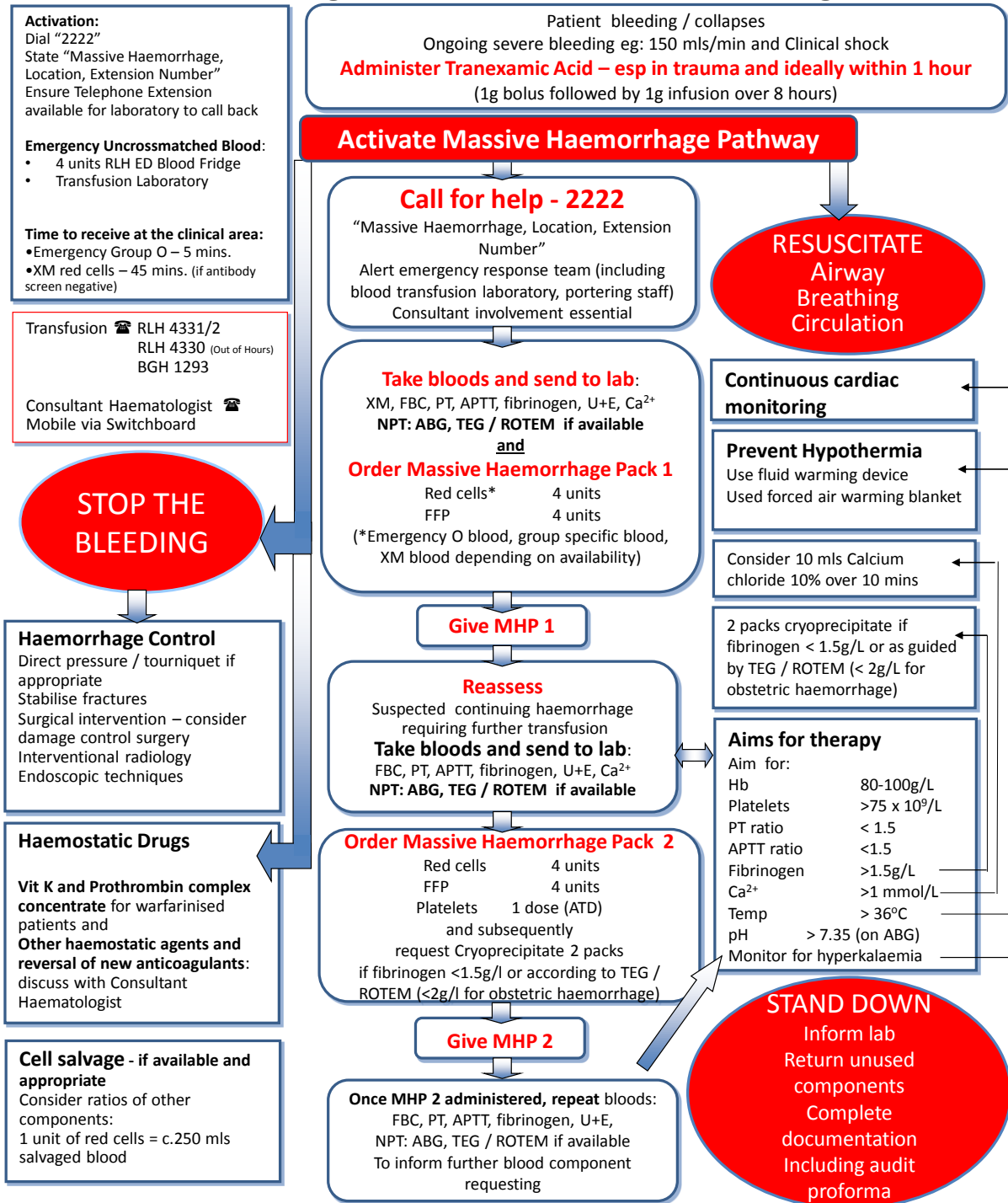
4.7 Stand down

Communication lead to communicate stand down of pathway to Laboratory and ensure used components have been fated on electronic blood tracking system. Any unused components to be returned to Laboratory in a timely manner.

4.8 Ensure documentation is complete

- Clinical area: monitoring of vital signs, timings of blood samples and communications, transfusion documentation in patient record, blood components fated on blood tracking.
- Laboratory: keep record of massive transfusion form

Transfusion Management of Massive Haemorrhage in Adults



Thromboprophylaxis should be considered when patient stable

ABG – Arterial Blood Gas
FFP- Fresh Frozen plasma
PT- Prothrombin Time

APTT – Activated partial thromboplastin time
MHP – Massive Haemorrhage Pack
TEG/ROTEM- Thromboelastography

ATD- Adult Therapeutic Dose
NPT – Near Patient Testing
XM - Crossmatch

Laboratory Management of Massive Haemorrhage

Massive Haemorrhage Pathway Activated

Transfusion receives Call
 'Massive Haemorrhage, Location, Specialty'
 Transfusion lab to liaise with other labs if appropriate

Receive call from designated communication lead in clinical area:
 'This relates to massive haemorrhage situation'
 The caller will state: Lab to complete **Massive Haemorrhage Form part 1**
Part 2 clinical area form to be sent with first blood components
 Name and contact telephone number, name of consultant responsible
 • Patient's ID (surname, forename, hospital number, DOB or minimum acceptable patient identifiers if unknown)
 • Requirements:
 • Whether O Neg/Pos is to be used
 • Order massive haemorrhage pack 1 +/- other components deemed appropriate.
 • Clarify urgency of requirements to decide on need for further emergency group O , or time to wait for group specific or crossmatched red cells (issue as part of pack 1)
 • FBC, PT, APTT, Fibrinogen, U+E*, ABG*, Calcium*, lactate*
 *may be near patient test or communication lead to liaise with appropriate labs

Receive samples and request forms

Haematology
 Perform FBC, PT, APTT, Fibrinogen

Transfusion
 Perform Group, antibody screen and crossmatch
Prepare MHP 1
 Red cells* 4 units
 (*emergency group O blood, group specific blood, XM'd blood depending on urgency)
 FFP (group specific) 4 units
 (If no plt stock order 2ATD on blue light from NHSBT)

Ring results to communication lead when available

Ring clinical area (communication lead) when blood / components ready

Receive further calls from communication lead in clinical area:
 Repeat investigations
 Order for MHP 2
 Liaise with on call haematologist (consultant / SpR) if needed
 Order for further components dependent on ongoing results
 Stand down

Prepare MHP 2
 Red cells 4 units
 FFP 4 units
 Platelets 1 ATD
 Cryoprecipitate 2 units if Fibrinogen < 1.5g/l

Transfusion Management of massive haemorrhage in children

Ensure a consultant is aware of the massive haemorrhage and a senior member of staff is available to take charge of resuscitation if not already present

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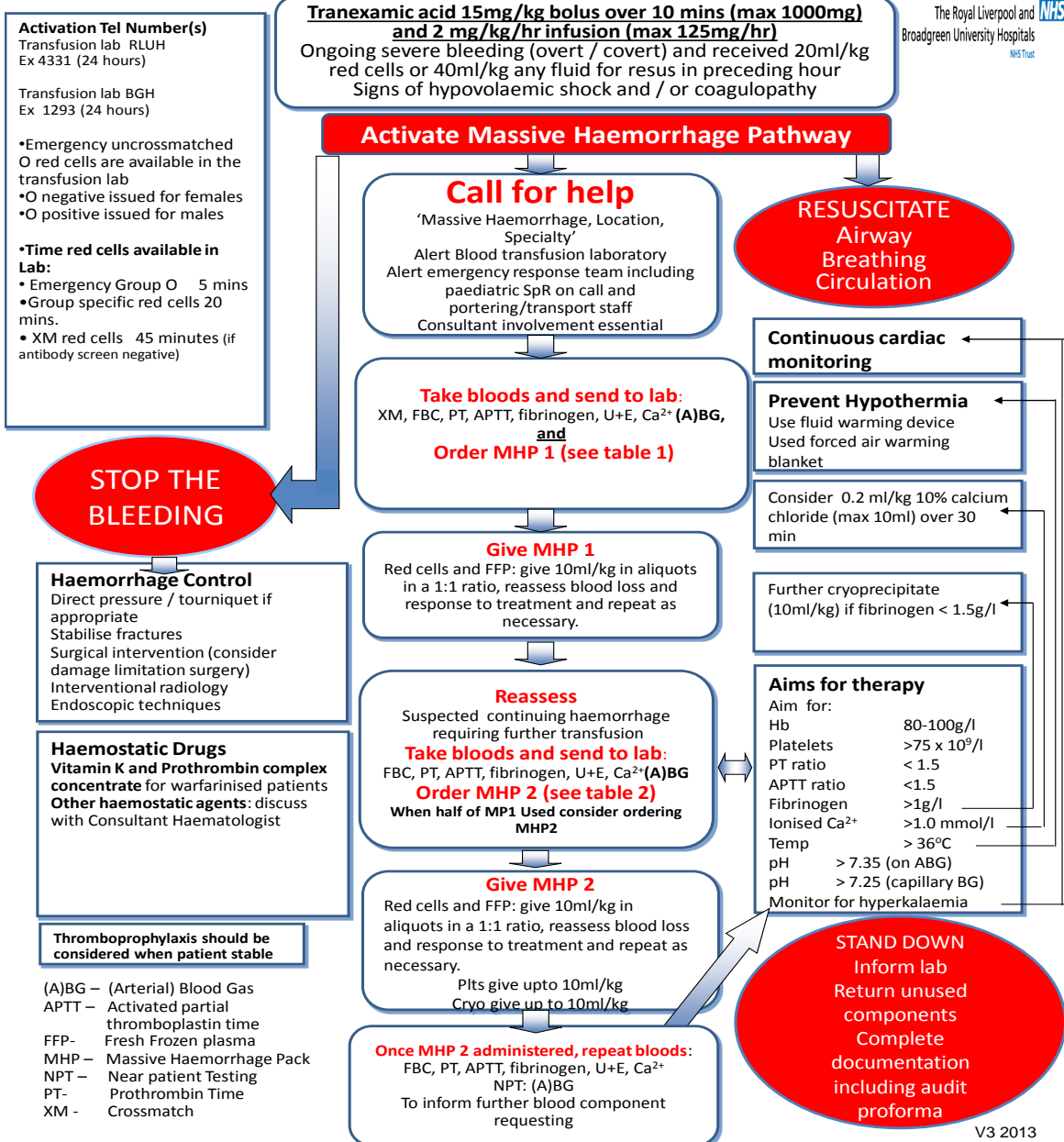


Table 1 – Major Haemorrhage pack 1 (MHP 1) order these volumes, which are also the maximum volumes to be administered from this pack in each weight category. Calculate volumes to be administered as detailed in the flow chart, but do not exceed these maximums (see example below)

Weight	Red cells * group O, group specific, crossmatched depending on availability	FFP
<5kg	2 Paediatric units (80-100ml)	100ml of OCTAPLAS
5-10kg	1 Adult unit (suitable for neonatal use if less than 12 months old) (250ml)	1 unit of OCTAPLAS
10-20kg	2 Adult units (suitable for neonatal use if less than 12 months old) (500ml)	2 units of OCTAPLAS
>20kg	4 Adult units (1000ml)	4units of OCTAPLAS

Table 2 – Major Haemorrhage pack 2 (MHP 2) order these volumes, which are also the maximum volumes to be administered from this pack in each weight category. Calculate volumes to be administered as detailed in the flow chart, but do not exceed these maximums (see example below)

Weight	Red cells	FFP	Cryoprecipitate	Platelets
<5kg	2 Paediatric units (80-100ml)	100ml of OCTAPLAS	1 single donor unit MB treated (40ml)	1 Paediatric pack of platelets (50ml)
5-10kg	1 Adult unit (suitable for neonatal use if less than 12 months old) (250ml)	1 unit of OCTAPLAS	2 single MB donor units (80ml)	2 Paediatric packs of platelets (100ml)
10-20kg	2 Adult units (suitable for neonatal use if less than 12 months old) (500ml)	2 units of OCTAPLAS	1 pool (5 units) (200ml) not MB treated	1 Adult apheresis pack (200ml)
>20kg	4 Adult units (1000ml)	4units of OCTAPLAS	2 pools (10 units) (400ml) not MB treated	1 Adult apheresis pack (200ml)

Red cells and FFP may be given through the same cannula via a Y-connector or 3-way tap provided the connection to the cannula is a short line. Platelets are ideally infused through a separate line, or after a clear flush, but may be given infused with red cells or FFP at a Y-connector or 3-way tap with a short connection to the cannula, **but the mixing must only occur after the platelets have passed through the filter.**

Administer red cells and FFP in aliquots of 10 ml/kg and in a ratio of 1:1; constantly assessing and reassessing the extent and rate of blood loss and the response to each such aliquot.

When half of MHP1 has been administered consider ordering MHP 2, if bleeding is on-going and control of the situation remains elusive.

Continue to administer aliquots of red cells and FFP in 10 ml/kg boluses as dictated by the patient's response to fluids, rate of blood loss etc (the whole clinical picture) until MHP2 is available.

With MHP2 use Red cells and FFP in the same fashion and administer a dose of platelets via a separate line (if at all possible) and give up to 10 ml/ kg of platelets. In addition administer a bolus of cryoprecipitate in a dose of up to 10 ml/kg .

Stop administering red cells and FFP if the patient's condition stabilises and it does not seem to be clinically indicated. Fine tune what products to give and in what volumes based on the lab results (when these become available) or TEG / ROTEM and bedside evidence of coagulopathy (microvascular bleeding).

Appendix 1 Document History and Version Control

Version	Date	Comments	Author
1.0	25 TH May 2018	Approved as part of the Blood Transfusion Policy.	Dr David Simcox
2.0	21/11/2018	Updated with new contact details to activate Massive Haemorrhage Protocol	Dr David Simcox

Review Process Prior to Ratification:

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