

MASSIVE TRANSFUSION PROTOCOL (MTP) - PAEDIATRIC PROCEDURE [®]

DOCUMENT SUMMARY/KEY POINTS

- Massive transfusion is defined as the replacement (or the anticipation of replacement) of >1 blood volumes within the first 24 hours of resuscitation, or >50% of blood volume within 3 hours.
- This procedure may be activated by any medical or nurse clinician when massive blood loss in a child is occurring or anticipated. The on call paediatric Haematologist will be involved to aid clinicians in management.
- There is mandatory minimum data set required by Blood Bank on activation
- Blood bank will dispense both red cell and plasma products in a 1:1 ratio.
- Frequent monitoring of venous blood gas, electrolytes and calcium as indicated.
- Randwick staff should adhere to the [SCH Flowchart for MTP](#)
- Westmead staff should adhere to the [CHW Flowchart for MTP](#)
- Westmead staff should use the [CHW Massive Blood Transfusion table](#)

Administration of all blood products must comply with **Transfusion of Blood and Blood Components policies:**

- **At CHW:** <http://webapps.schn.health.nsw.gov.au/epolicy/policy/1214>
- **At SCH:** <http://webapps.schn.health.nsw.gov.au/epolicy/policy/3597>

This document reflects what is currently regarded as safe practice. However, as in any clinical situation, there may be factors which cannot be covered by a single set of guidelines. This document does not replace the need for the application of clinical judgement to each individual presentation.

Approved by:	Policy, Procedure and Guideline Committee	
Date Effective:	1 st March, 2018	Review Period: 3 years
Team Leader:	Staff Specialist – SCH and CHW	Area/Dept: Emergency/Haematology

CHANGE SUMMARY

- Development of a Network Massive Transfusion Protocol: It replaces the site-based documents.
- Development of two separate flowcharts to reflect what occurs at both Hospitals.
- Site-specific flowcharts (with more site specific details) are linked documents that are printable for easy access.

READ ACKNOWLEDGEMENT

- Appropriate staff working in acute care areas and Haematology staff are to read and acknowledge they understand the contents of this document.

This document reflects what is currently regarded as safe practice. However, as in any clinical situation, there may be factors which cannot be covered by a single set of guidelines. This document does not replace the need for the application of clinical judgement to each individual presentation.

Approved by:	Policy, Procedure and Guideline Committee	
Date Effective:	1 st March, 2018	Review Period: 3 years
Team Leader:	Staff Specialist – SCH and CHW	Area/Dept: Emergency/Haematology

TABLE OF CONTENTS

Purpose	4
Definition	4
Activating Massive Transfusion Protocol (MTP)	4
<i>Medical or nursing staff may activate the MTP if:</i>	4
<i>Blood Bank may activate the MTP if:</i>	4
Use of intravenous tranexamic acid (Cyclokapron) in trauma	5
Massive Transfusion Pack	6
Immediate Dispatch.....	6
Second Dispatch	6
Time for Blood Product Availability	7
Laboratory Criteria – target values	8
Ongoing Clinical and Laboratory Assessment	8
Deactivation of MTP	8
Flowchart – Paediatric Massive Transfusion Protocol	9
Randwick Paediatric MTP Flowchart.....	9
Westmead Paediatric MTP Flowchart	9
Abbreviations	9

Purpose

This document describes the **activation and deactivation procedure** to provide blood components to patients requiring massive blood transfusion in a timely manner¹.

Definition

Massive transfusion is defined as the transfusion (or anticipation of transfusion) of

- one or more blood volumes within 24 hours, **OR**
- 50% of blood volume within 4 hours, **OR**
- 40mL blood/kg².

Activating Massive Transfusion Protocol (MTP)

Medical or nursing staff may activate the MTP if:

- Any child requiring more than 20mL/kg of packed red blood cells (PRBC) in 2 hours and anticipated ongoing blood loss;

OR

- Any child requiring more than 40mL/kg of PRBC in a 24 hour period with ongoing blood loss;

OR if at Westmead

- “Code Crimson” for trauma with acute life threatening haemorrhage is activated (see CHW Procedure: [Trauma: Code Crimson](#)).

Blood Bank may activate the MTP if:

- > 2 units of PRBC issued within 1 hour for child < 5 years old, **OR**
- > 4 units PRBC issued within 1 hour for child ≥ 5 years old, **OR**
- The Blood Bank technician anticipates likelihood of additional component needs. In this situation, the Blood Bank technician contacts the Paediatric Haematologist to activate the protocol, **OR**
- If at Westmead*, “Code Crimson” is activated.

Note: An exception to the above criteria is planned cardiac bypass surgery. If blood loss or usage is greater than anticipated, the MTP may be initiated at the discretion of the surgeon or anaesthetist.

The massive transfusion protocol is activated by notifying the Blood Bank technician on-call.

At CHW, Westmead: call ext. 52284 or pager 6832
At SCH, Randwick: call ext. 29145

The Blood Bank technician will then notify the on-call Paediatric Haematologist, who will liaise with the clinical team.

Details required by Blood Bank

When providing (verbal) information to Blood Bank, the following details are required:

- **Name and contact (phone/pager) of contact person.**
(It is best to identify one person who will co-ordinate with Blood Bank³. e.g. anaesthetist in Operating Suite, surgeon or emergency physician in the Emergency Department.)
- **Name and medical record number of patient.**
- **Weight of patient.**
- **Location of patient and phone number of location.**
Note: Clinical team must notify Blood Bank if the patient location changes.
- **Initial blood results**
e.g. full blood count (FBC), coagulation screen (coags), venous blood gas (VBG), cross match.
- **Blood components transfused prior to arrival at hospital/activation of the Massive Transfusion protocol.**
- **Urgency of need for blood products**
e.g. immediate or over next 30 minutes³. This will help determine what products will be despatched. Refer to [Time for Blood Product Availability table](#).

Use of intravenous tranexamic acid (Cyclokapron) in trauma

The use of intravenous tranexamic acid in trauma has been established for adult patients (CRASH-2 trial)⁴ and administration is likely to be beneficial in children as well:

- Tranexamic acid at a dose of 15 mg/kg (dose range 10-20 mg/kg; max 1g) by slow intravenous infusion over 10-15 minutes; up to the first 3 hours after trauma.
- Follow up doses can be given at the decision of the treating team and can be as IV boluses or continuous infusion
- Precautions:
 - Avoid rapid infusion or push injection as it may cause hypotension.
 - Tranexamic acid should be used with caution in patients with haematuria, renal haemorrhage and bleeding into other bodies cavities (e.g. pleural space, joints) as inhibition of fibrinolysis may result in retention of blood clots in those spaces.
 - Dose/interval adjustments for subsequent doses are required in renal impairment
 - Do not use in conjunction with factor IX complex concentrates

Massive Transfusion Pack

Immediate Dispatch

The following “pack” will be despatched immediately upon activation of the MTP⁵ according to patient weight:

Weight of Child			
< 15kg	15 – 30kg	30 – 50kg	> 50kg
1 unit PRBC ⁺	2 unit PRBC	3 unit PRBC	4 unit PRBC
1 unit FFP [*]	2 unit FFP	3 unit FFP	4 unit FFP
2 units cryoprecipitate	3 units cryoprecipitate	5 units cryoprecipitate	8 units cryoprecipitate

Specimen tubes for sample collection from patient after administration of products will be sent with each pack. Pathology form (with required tests pre-printed) will be included.

+ PRBC = Packed Red Blood Cells

* FFP = fresh frozen plasma

Second Dispatch

Upon request for a second lot of blood products, the following will be issued according to weight:

Weight of Child			
< 15kg	15 – 30kg	30 – 50kg	> 50kg
1 unit PRBC	2 unit PRBC	3 unit PRBC	4 unit PRBC
1 unit FFP	2 unit FFP	3 unit FFP	4 unit FFP
1 unit pooled platelets	1 unit pooled platelets	1 unit pooled platelets	1 unit pooled platelets

Specimen tubes for sample collection from patient after administration of products will be sent with each pack. Pathology form (with required tests pre-printed) will be included.

By this stage, the on-call paediatric Haematologist will be involved, and will direct further products in consultation with the clinical team. The principles of further replacement will be:

- 1:1 ratio of PRBC to FFP^{5 - 12},
- Alternating platelets and cryoprecipitate, but adjusted according to laboratory results³.
- Factor concentrates such as activated factor VII (“Novoseven”) or prothrombin complex concentrates (“Prothrombinex”) may be indicated, and this will be the decision of the on-call paediatric Haematologist in consultation with the clinical team^{3, 13-15}.

Note: Administration of all blood products *must comply* with the local Transfusion of Blood and Blood Components policy.

Responsibilities

The following points outline the responsibilities throughout the activation of the massive transfusion protocol (MTP) for Blood Bank and the clinical service/team involved.

Blood Bank Service

- Prioritise testing and product distribution to patient.
- Despatch massive transfusion package to the appropriate location.
- Maintain communication with clinical team and ensure continuous availability of products. Commence thawing of next batch of frozen products immediately upon despatch of previous pack.
- Notifies and communicates with on-call paediatric Haematologist.

Clinical Service

- Ensure availability of runner to transport products rapidly to patient location.
- Informs Blood Bank as soon as possible if patient location changes.
- Deactivates MTP by notifying Blood Bank.

Time for Blood Product Availability

Product	Time Until Despatch From Blood Bank
O negative PRBC	Immediate
ABO specific PRBC (uncrossmatched)	5 – 10 minutes
Crossmatched PRBC	40 minutes
Frozen products (FFP, cryoprecipitate)	20 – 30 minutes

Laboratory Criteria – target values

As a general guideline in massive transfusion, the following target values are reasonable to aim for:

Test	Target value
Haemoglobin (Hb)	> 70 g/L
Platelets	> 50 x 10 ⁹ /L (or > 100 x 10 ⁹ /L if neurological injury) ^{3, 5, 7}
Activated partial thromboplastin time (APTT)	< 40 seconds ⁴
Prothrombin time (PT)	< 20 seconds
Fibrinogen	> 1 g/L ^{3, 7}
Acidosis and hypothermia should also be vigorously corrected, as these exacerbate coagulopathy ^{3, 5, 13, 16} (refer to Ongoing Clinical and Laboratory Assessment).	

Ongoing Clinical and Laboratory Assessment

The following points need to be actioned/considered:

- Strict compliance with identification, documentation and product administration is mandatory as per local Transfusion of Blood and Blood Components policy.
- Laboratory tests (FBC, coags) **after administration of each “pack”**.
- Additional crossmatch samples may be required. Clinical team will be notified by Blood Bank technician.
- Frequent monitoring of venous blood gas, electrolytes and calcium as indicated.
- Consider using a blood warmer.
- Consider use of cell saver (available in Operating Suite) to scavenge blood or to wash blood products and also rapid transfuser in children >30kgs. Inform theatre floor manager as early as possible if you require cell saver as cardiac technicians need to be called in to set this up. Allow about 30 minutes for this to happen.

Deactivation of MTP

When deactivating the MTP the following will occur:

- The clinician in charge of the patient (or delegate) will notify Blood Bank of decreasing needs for products or termination of the massive transfusion protocol.
- Blood Bank technician will notify on-call paediatric Haematologist.

Flowchart – Paediatric Massive Transfusion Protocol

Randwick Paediatric MTP Flowchart

- <http://webapps.schn.health.nsw.gov.au/epolicy/my/requests/3975/attachments/3967/download>

Westmead Paediatric MTP Flowchart

- <http://webapps.schn.health.nsw.gov.au/epolicy/my/requests/3975/attachments/3968/download>

Abbreviations

MTP	Massive Transfusion Protocol
PRBC	Packed red blood cells
FBC	Full blood count
Coags	Coagulation Screen
VBG	Venous blood gas
APTT	Activated partial thromboplastin time
PT	Prothrombin time
FFP	Fresh frozen Plasma

Copyright notice and disclaimer:

The use of this document outside Sydney Children's Hospitals Network (SCHN), or its reproduction in whole or in part, is subject to acknowledgement that it is the property of SCHN. SCHN has done everything practicable to make this document accurate, up-to-date and in accordance with accepted legislation and standards at the date of publication. SCHN is not responsible for consequences arising from the use of this document outside SCHN. A current version of this document is only available electronically from the Hospitals. If this document is printed, it is only valid to the date of printing.