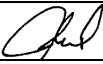




**Document History:**

**Title:** Receiving Blood, Blood Components and Plasma Protein Products (Derivatives)      **Site(s):** All DSM sites

<b>Document #:</b>	<b>160-INV-07</b>	<b>Version #:</b>	<b>04</b>
<b>Section:</b>	<b>Manitoba Transfusion Quality Manual for Blood Banks</b>	<b>Subsection:</b>	<b>INV Module</b>

<b>Approved by:</b>	<u>C Musuka</u>	<b>Written By:</b>	<u>TM Discipline Team</u>
Signature:			
Date:	<u>31-JAN-2017</u>	Date:	<u>March 2011</u>

#	Details of Revisions:	Approval:	Date:
1	New document	A Kabani	31-MAR-2011
2	<ul style="list-style-type: none"> <li>Added to 4.0, 5.0, 6.0 for Trace Line sites to refer to Trace Line Procedure</li> <li>Revised example of receipt label to include enter in Trace Line</li> <li>Revised example of receipt label to "or Inter Facility Transfer form" not "or INV.002"</li> </ul>	C Musuka	05-NOV-2012
3	<ul style="list-style-type: none"> <li>2.4 New scope to notify ward if applicable</li> <li>Ward notified updated on label</li> </ul>	C Musuka	18-APR-2013
4	<ul style="list-style-type: none"> <li>2.5 to 2.9 new</li> <li>3.0 Materials added TL packing slip and removed CBS feedback form</li> <li>4.0, 5.0 and 6.0 revised Procedures for TL sites</li> <li>Revised throughout to update to PPP</li> </ul>	C Musuka	31-Jan-2017
5			
6			
7			

## Receiving Blood, Blood Components and Plasma Protein Products (Derivatives)

### 1.0 Principle

- 1.1 To receive blood, blood components and plasma protein products into inventory.
  - from the blood supplier, via inter-facility transfer or shipped with a patient
- 1.2 To provide an accurate record of the receipt of blood, blood components, and plasma protein products

### 2.0 Scope and Related Policies

- 2.1 Blood, blood components and plasma protein products shall be handled, stored, distributed, and shipped in a manner that prevents damage and limits deterioration.
- 2.2 A process shall be in place to ensure the traceability of all blood, blood components and plasma protein products received.
- 2.3 A process shall be in place to ensure that segments from all transfused units are removed and stored at 1°C to 6°C for a minimum of 7 days after transfusion.
- 2.4 A process shall be in place to notify clinical ward upon receipt of any patient specific blood, blood components or plasma protein products if applicable
- 2.5 To promote traceability, designated Trace Line Testing or Hub sites have been established as site for Non-Trace sites to receive blood products from
  - site is dependent on; Health region and established transportation routes
- 2.6 Red cells, thawed plasma and platelets received from DSM Trace Line testing site/Hub site to non-Trace Line facility shall have data logger enclosed
  - including stock emergency red cells and patient specific units
- 2.7 Upon receipt of shipment with data logger enclosed the receiving facility shall:
  - Remove temperature data logger from box and record date and time of unpacking on shipping container tracking form (F160-QCFORM-38)
  - Retrieve data from data logger and verify shipping temperature during shipment
  - Save data in appropriate file on computer at facility
  - Take temperature upon receipt using calibrated thermometer if; data logger did not record data, site unable to download date or temperature during shipment was outside required limits
  - Return data logger and Shipping Container Tracking Form (F160-QCFORM-38) to shipping facility
- 2.8 Contact shipper, and complete customer feedback form, and if applicable TM Physician consult initiated if:
  - There is no security seal on the shipping box
  - Transport time exceeded 24 hours
  - Discrepancies with packing slip
  - Packing errors
  - For shipments with data loggers; unacceptable temperature during shipment, data logger did not record any data, or site unable to download data

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**2.0 Scope and Related Policies cont'd**

- 2.9** The packing configuration for blood, blood component and plasma protein products varies and is dependent if product received from CBS or DSM site and shipping containers used
- Frozen blood components shall only be received from CBS
  - For acceptable packing configuration for product received from DSM sites refer to INV Procedure: Inter-Facility shipping of Blood, Blood Components and Plasma Protein Products (Derivatives)
  - For CBS packing configuration. Refer to CBS customer letter 2017-03 at : <https://blood.ca/en/hospital/customer-letters>

**3.0 Materials**

- CBS Packing slip(s)
- Trace Line Packing Slip
- Blood, Blood Component, Plasma Protein Product (Derivative)
- Shipping container with appropriate packing materials
- Record of Transfusion (ROT)
- BTS/CBS Patient Report
- Appropriate blood bank log
- INV Form- Inter-Facility Blood, Blood Component and Derivative Transfer
- Product tag(s)
- INV Form- DSM Blood Bank Customer Feedback Form
- Thermometer (for receiving blood and thawed components)
- TH10 Temperature USB Datalogger
- Shipping Container Tracking Form (F160-QCFORM-38)
- Receipt Label

**Example of Receipt Label:**

Received From _____		Received Date and Time _____	
Temperature on Receipt (blood only) _____		Travel Time _____	
<input type="checkbox"/> N/A			
Received By _____			
<u>Security Seal Intact</u>		<u>Segments removed</u>	
<input type="checkbox"/> Yes		<input type="checkbox"/> Yes	
<input type="checkbox"/> No		<input type="checkbox"/> No	
<u>Packaging</u>		<u>ABO/Rh Confirmation Testing Red Cell Units</u>	
<input type="checkbox"/> Acceptable		<input type="checkbox"/> Yes	
<input type="checkbox"/> Unacceptable		<input type="checkbox"/> N/A	
<u>Product check with packing slip or Inter Facility Transfer Form</u>			
<input type="checkbox"/> Yes <input type="checkbox"/> No			
<u>Visually inspected and found suitable</u> <input type="checkbox"/> Yes <input type="checkbox"/> No			
<u>Ward Notified</u> <input type="checkbox"/> Yes <input type="checkbox"/> NA			
Entered in appropriate blood bank log/Trace Line <input type="checkbox"/>			

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### 4.0 Procedure

For Trace Line Sites refer to Trace Line Procedures:

- Receipt of Blood, Blood Components and Plasma Protein Products (from CBS) in Trace Line
- Handling in Trace Line: Receipt of Inter-Facility Shipment of Blood, Blood Components and Derivatives

For Blood see INV Procedure- Receipt of Blood

For Blood components see INV Procedure- Receipt of Blood Components

For Derivatives see INV Procedure- Receipt of Plasma Protein Products (Derivatives)

### 5.0 Reporting

For Trace Line Sites refer to Trace Line Procedures:

- Receipt of Blood, Blood Components and Plasma Protein Products- Derivatives (from CBS) in Trace Line
- Handling in Trace Line: Receipt of Inter-Facility Shipment of Blood, Blood Components and Derivatives

For Blood see INV Procedure- Receipt of Blood

For Blood components see INV Procedure- Receipt of Blood Components

For Derivatives see INV Procedure- Receipt of Plasma Protein Products (Derivatives)

### 6.0 Procedural Notes

For Trace Line Sites refer to Trace Line Procedures:

- Receipt of Blood, Blood Components and Plasma Protein Products- Derivatives (from CBS) in Trace Line
- Handling in Trace Line: Receipt of Inter-Facility Shipment of Blood, Blood Components and Derivatives

For Blood see INV Procedure- Receipt of Blood

For Blood components see INV Procedure- Receipt of Blood Components

For Derivatives see INV Procedure- Receipt of Plasma Protein Products (Derivatives)