



Blood Products  
Stem Cells  
Human Tissues



Hema-Quebec  
Stem Cell Donor Registry  
4045 Cote-Vertu, St-Laurent  
QC, Canada, H4R 2W7  
Tel : + 514-832-1031  
Fax : + 514-832-0266  
www.hema-quebec.qc.ca

## COLLECTION CENTRE UNSTIMULATED LEUKAPHERESIS PRODUCT (MNC, APHERESIS) REPORT

*(To be completed by collection centre to confirm collection details and product analysis)*

Hema-Quebec Recipient ID#	Hema-Quebec Donor ID#:
International Recipient ID#:	Donor DOB (dd/mm/yyyy):
Collection Centre:	Donor ABO/Rh:
Date of Collection (dd/mm/yyyy):	Donor Gender: <input type="checkbox"/> M <input type="checkbox"/> F

**PRODUCT ANALYSIS FOR DAY 1 COLLECTION: Collection Date: (dd/mm/yyyy):** \_\_\_\_\_

<b>Volume of Blood processed:</b> _____	
WBC: _____ x 10 <sup>9</sup> /L x Total volume of product: _____ mL = Total Nucleated Cells: _____ x 10 <sup>9</sup>	
MNC: _____ x 10 <sup>9</sup> /L x Total volume of product: _____ mL = Total Nucleated Cells: _____ x 10 <sup>9</sup>	
Total CD 3+ cells collected: _____ x 10 <sup>8</sup>	
Document volume and concentration of any media, anticoagulant and antibiotics that were added to the product post collection: Record lot number and expiry date if applicable	
Additive: _____ ml Conc. _____ Lot No: _____ Expiry Date: _____	
Additive: _____ ml Conc. _____ Lot No: _____ Expiry Date: _____	
Product Storage: Time _____ Time Zone: _____ Temperature _____ (if applicable)	
Any changes in collection requirements or comments?	Any changes in peripheral samples requested? If yes please specify:

Signature: \_\_\_\_\_ Date (dd/mm/yyyy): \_\_\_\_\_



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**PRODUCT ANALYSIS FOR DAY 2 COLLECTION: Collection Date: (dd/mm/yyyy):** \_\_\_\_\_

<b>Volume of Blood processed:</b> _____	
WBC: _____ x 10 <sup>9</sup> /L x Total volume of product: _____ mL = Total Nucleated Cells: _____ x 10 <sup>9</sup>	
MNC: _____ x 10 <sup>9</sup> /L x Total volume of product: _____ mL = Total Nucleated Cells: _____ x 10 <sup>9</sup>	
Total CD 3+ cells collected: _____ x 10 <sup>8</sup>	
Document volume and concentration of any media, anticoagulant and antibiotics that were added to the product post collection: Record lot number and expiry date if applicable	
Additive: _____ ml Conc. _____ Lot No: _____ Expiry Date: _____	
Additive: _____ ml Conc. _____ Lot No: _____ Expiry Date: _____	
Product Storage: Time _____ Time Zone: _____ Temperature: _____ (if applicable)	
Any changes in collection requirements or comments?	Any changes in peripheral samples requested? If yes please specify:

Signature: \_\_\_\_\_ Date (dd/mm/yyyy): \_\_\_\_\_

<b>ATTESTATION TO LABELLING REQUIREMENTS</b>	
Courier: I have examined all products, samples, and documents and verify that they are accurate and complete according to the patient and donor identifiers as indicated above.	
Name of Courier: _____	Signature of Courier: _____ Date (dd/mm/yyyy): _____
Product Received by Courier (dd/mm/yyyy) _____	Time Received: _____ Time Zone: _____
Collection Centre Representative: _____	Signature: _____
Date (dd/mm/yyyy) : _____	

*Courier- original; copy to Collection Centre and to Héma-Québec at +(514)832-0266.*



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## COLLECTION CENTRE UNSTIMULATED LEUKAPHERESIS PRODUCT (MNC, APHERESIS) REPORT

### NOTICE TO TRANSPLANT CENTRE

**By accepting the accompanying product, you (the transplant centre) agree to the following:**

- The cell products collected from this donor are intended solely for the purpose of immediate infusion for the above mentioned patient.
- Excess cells may be cryopreserved and stored for future therapeutic treatment for this patient. No other uses of these cells are permissible.
- Cells not used for the therapeutic treatment of the above mentioned patient must be properly discarded.
- Héma-Québec must be provided with detailed information concerning the use and/or disposal of this cellular product, whole or partial.
- Any requests deviating from these terms must be submitted in writing to Héma-Québec for approval prior to collection. Héma-Québec reserves the right to accept or refuse any of these requests. (ex. Cryopreservation of total product)

### ***To be completed by Transplant Centre:***

The above collection specifications are:  accepted  not accepted

Transplant Physician signature: \_\_\_\_\_

Date (dd/mm/yyyy): \_\_\_\_\_

Name of Transplant Physician (please print): \_\_\_\_\_