



**COLLECTION REPORT: STIMULATED HUMAN PERIPHERAL BLOOD STEM CELL
(HPC, Apheresis)**

(To be completed by the collection center)

Overnight storage method/details:
Transport Temperature: (Special packing materials such as gel packs must be provided by the transplant center unless alternative arrangements have been made with the donor or collection center)

MATERIAL:

Material	Commercial name	LOT number	Expiration date	Inspection	Total amount
ACD				<input type="checkbox"/> OK	ml
Heparin				<input type="checkbox"/> OK	
Collection kit				<input type="checkbox"/> OK	
				<input type="checkbox"/> OK	
				<input type="checkbox"/> OK	

CELL COUNT:

Volume collected:	ml			
NC COUNT	/ μ l		TOTAL:	10^8
CD 34 +:	%			
CD 34+:	/ μ l		TOTAL:	10^6
Calculated CD 34+ cell dose for recipient				
Prescribed		10^6 CD34+/ kg	=	10^6 CD34+/kg
Collected		10^6 CD34+/ kg	=	10^6 CD34+/kg

OTHER QUALITY CONTROL:

Peripheral blood WBC count: / μ l
Culture for bacterial contamination: <input type="checkbox"/> Negative / <input type="checkbox"/> Positive / <input type="checkbox"/> Not done
If positive: give details:
Other tests done:



Marrow Donor Program Belgium – Registry
Motstraat 40 2800 Mechelen
Tel: (32) - 15 44 33 96 Fax: (32) - 15 44 36 56
Email : MDPB-registry@rodekruis.be

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ADDITIONAL SAMPLES:

Type of sample		Prescribed	Collected
PBSC:		ml	ml
Blood:	CLOTTED (no anti-coagulant)	ml	ml
	EDTA	ml	ml
	HEPARIN	ml	ml
	ACD	ml	ml

DISCLAIMER: The cell products collected from this donor are intended solely for the purpose of immediate therapeutic treatment for the above-mentioned patient. Excess cells may be stored for future infusion 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 disposed of properly. The donor center must be provided detailed information concerning the use and/or disposal of all portions of this cell product. By accepting these cells, the transplant physician also accepts these terms and conditions. Requests for deviations from these terms must be submitted in writing to the donor center for approval.

PROBLEM REPORTED BY TRANSPLANT CENTER:

Problem reported:
Action taken:

Collection physician: STAMP:	Signature:	Date: (Day/Month/Year)
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