

# (archived) CBB Survey 2015 WO-1367

## CBB Survey 2015

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<b>1. General Information</b>	
Name of CBB	CBB Roma Azienda Policlinico Umberto I
CBB Director	Gabriella
CBB Director	Girelli
Address	Via Chieti, 7
Address	00161
Address	Rome
Phone Number	0039.06.49976549
Date CBB Started Collecting Cord Blood Units (month/day/year)	06/01/1994
Number of Public Cord Blood Units	1,185
Planned Number of Public Cord Blood Units Stored in 2015	100
Lists on BMDW	Yes
Affiliated with National Stem Cell Donor Registry	Yes
Registry Affiliation	IBMDR
<b>2. Cord Blood Units in Inventory</b>	
Current Processing Method	Liquid+ plasma only
Year Current Process Method Started	2009
Percent of Units Plasma and RBC Reduced (manual)	20
Percent of Units No Volume Reduction	80
<b>3. Accreditations, Licenses and Certifications</b>	
FACT-Netcord	No
AABB	No
Competent Authority/ National Health Authority	Yes
Name of Competent Authority	National Blood Center
Audited by National Stem Cell Registry	No
ISO	Yes
<b>4. Cord Blood Collection</b>	
Current Collection Practice Is the collection In/Ex -Utero or both?	Both
Current Antiseptic	Chlorhexidine
Current Antiseptic	Betadine
Collection Bag	Double needle
Agitation during Collection	Automatic

<b>5. Conditioning/Transport from Collection Site to CBB</b>	
Secondary Bag Used	Yes
Transport Conditions	Qualified transporter
Transport Conditions	Insulating transport container
Transport Conditions	Active refrigeration system
Transport Conditions	Electronic temperature probe
Transport Conditions	Ground transport
Temp. for Storage and Transport	Other (lower limit +1-35°C, higher limit +6-30°C)
<b>6. Pre-Processing Evaluation</b>	
Completed Prior to Accepting a CBU	Medical history, collection report, informed consent
Method for CD34 Remuneration	ISHAGE guidelines
External Proficiency Testing for QC of FACS Lab	Yes
External Proficiency Testing for QC of FACS Lab	Missing
Post Processing/ Pre Freeze CD34+ Cell Count	Yes
Time from Collection to Processing	up to 48H
<b>7. Processing and Packaging</b>	
Pre Freeze Processing Methods- Unit in Inventory	No processing
Pre Freeze Processing Methods- Unit in Inventory	Other: Manual plasma depletion
Pre Freeze Processing Methods- Current	Other: Plasma depletion manual
Additives Currently in Use	No additive
Current Cryopreservation Method	Conventional CRF
Current Cryoprotectant Additive	DMSO
Current Cryobag	Single bag (one fraction)
Current Target Cryopreservation Volume (mL)	bag ml : 10-30 ; 30-60; 60
Current Packaging for Storage	Canister
Current Packaging for Storage	More than one segment
Current Packaging for Storage	Other: Not attached vials stored with CBU
<b>8. Testing</b>	
PCR Performed on IgM+ Result	CMV
PCR Performed on IgM+ Result	Toxoplasmosis
PCR Performed on IgM+ Result	EBV
Extra Material Currently Stored	Cord blood DNA
Extra Material Currently Stored	Plasma/cord blood
Extra Material Currently Stored	Maternal DNA
Extra Material Currently Stored	Maternal plasma/serum
Current Post Processing Threshold for Accepting a CBU for Public Use TNC	120
Current Post Processing Threshold for Accepting a CBU for Public Use CD34 (10^6) Single Platform	1.00
Current Post Processing Threshold for Accepting a CBU for Public Use CD34 (10^6) Double Platform	NA

Current Post Processing Threshold for Accepting a CBU for Public Use CFU-GM	Growth
Current Post Processing Threshold for Accepting a CBU for Public Use CFU	Growth
Current Post Processing Threshold for Accepting a CBU for Public Use Viability	85
<b>9. Storage</b>	
Type of Storage Container Used	Conventional tank liquid phase
Monitoring of Storage	Centralized alarm system local
<b>10. HLA Typing</b>	
Current Level of HLA Typing at Time of Listing HLA-A	LR
Current Level of HLA Typing at Time of Listing HLA-B	LR
Current Level of HLA Typing at Time of Listing HLA-C	
Current Level of HLA Typing at Time of Listing HLA-DRB1	HR
Current Level of HLA Typing at Time of Listing HLA-DQB1	
Current Level of HLA Typing at Time of Listing HLA-DPB1	
Accreditation of HLA Lab	EFI accredited lab
Average Turnaround Time for Extended HLA Typing Results in days	?10
Attached Segment Used for Confirmatory/ Verification Typing	Yes
Units Listed without Attached Segment and have not been Previously Typed on Attached Segment	Yes
Percentage of CBUs that have an Attached Segment	75-90%
Confirmatory/ Verification Typing on an Attached Segment is Pre-Release Requirement	Yes
<b>11. Reservation and Cancellation Policies</b>	
What Point is a CBU Reserved for a Patient	Reservation request
Length of Time a CBU can be Reserved in days	60
Reservation Fee	No
Reservation Cancellation Fee in Absence of Shipment Request	No
Can Reservation be Extended	Yes
Is a Unit Report Provided on a Unit that is Reserved for Another Patient	No
Is TC Informed when CBU is Released	Yes
<b>12. Release and Shipment</b>	
Hemoglobinopathy Screening Performed Prior to Release	Yes
Criteria to Ship a CBU Viability and Cell Count	50
Criteria to Ship a CBU HLA Identity Testing	Yes
Current Packaging for Shipment to TC	Metal canister
Current Packaging for Shipment to TC	One attached segment

Current Packaging for Shipment to TC	Separate segment
Time Between Shipment Request and Sending CBU	1-2 weeks
Fee for Shipment Cancellation	No
Dry Shippers Validated to Maintain Temperature of at least -150 for 48 hours Beyond Expected Arrival	Yes
Electronic Temperature Data Logger on All Dry Shippers	Yes
Who Selects Transport Company	CBB
Who Selects Transport Company	world courier selected by IBMDR
Shape of Transport Container	Mushroom
<b>13. Adverse Events Reporting</b>	
Who are S(P)EARS Reported To Competent Authority	Yes

14. Pictures of cord blood units in the inventory

15. Infectious Disease Marker (IDM) CURRENTLY performed.




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## Holiday Calendar

Team Calendars
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