

# CB banking specifics WO-1886

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## Section 1 General Info

| Question   | CBB answer |
|--|------------|
| Name of the cord blood bank:   |            |
| Number of cord blood units the cord blood bank plans to store for public use (unrelated patients): |            |

## Section 2 Cord Blood Units in Inventory

| Question  | CBB answer |
|---|------------|
| Current processing method(s):   |            |
| Plasma and RBC reduced (automatic)  | checkfalse |
| Plasma reduction only   | checkfalse |
| Plasma and RBC reduced (manual)   | checkfalse |
| RBC reduction only  | checkfalse |
| Total Nucleated Cell Count (x10E7) of your cord blood units stored for Unrelated Patients (Public Use). |            |
| < 125 :   |            |
| 125 - 149 :   |            |
| 150 - 199 :   |            |
| 200 - 250 :   |            |
| > 250 :   |            |

## Section 3 Cord Blood Collection

| Question                                    | CBB answer |
|---|------------|
| Current practice for collecting cord blood: |            |
| Current antiseptic:                         |            |
| Collection bag:                             |            |
| Agitation during collection:                |            |

## Section 4 Conditioning and transport from Collection Centre to cord blood bank

| Question   | CBB answer |
|--|------------|
| Secondary bag used by the cord blood bank (sealed, plastic bag or similar to avoid any leakage): |            |
| Transport conditions:  |            |
| Insulating transport container   | checkfalse |
| Active refrigeration system  | checkfalse |
| Passive refrigeration system (gel, blocks)   | checkfalse |
| Electronic temperature probe   | checkfalse |
| Non-electronic temperature probe   | checkfalse |
| Qualified transporter  | checkfalse |
| Unqualified transporter  | checkfalse |
| Air transport  | checkfalse |
| Ground transport   | checkfalse |
| Other,   | checkfalse |
| Temperature range for storage and transportation of fresh product:                               |            |

## Section 5 Evaluation

| Question   | CBB answer |
|--|------------|
| Pre-processing Evaluation:<br>Current threshold for accepting a cord blood unit for public use in the cord blood bank: |            |
| Net weight in grams (excluding bag and anticoagulant) before processing  |            |
| TNC (10E7) before processing   |            |
| Viability CD45 positive cells (%)  |            |
| Viability CD45 positive cells (method)   |            |
| Viability CD34 positive cells (%)  |            |
| Viability CD34 positive cells (method)   |            |
| Criteria that are completed before accepting a cord blood unit for public use in the cord blood bank:                  |            |
| Medical History  | checkfalse |
| Collection Report  | checkfalse |
| Informed Consent   | checkfalse |
| Maternal IDM results   | checkfalse |
| Temperature and integrity of the bag   | checkfalse |
| Other,   | checkfalse |
| Used method for CD34 enumeration:  |            |
| The cord blood banks uses external proficiency testing for QC of the cord blood banks FACS lab:                        |            |
| The cord blood bank performs post processing/pre-freeze CD34 cell count:   |            |
| Acceptable time from collection to processing:   |            |

## Section 6 Processing-Packaging

| Question   | CBB answer |
|--|------------|
| The pre-freeze processing method(s) cord blood bank uses at any stage in time:       |            |
| AXP  | checkfalse |
| SEPAX  | checkfalse |
| Optipress  | checkfalse |
| Prepacyte  | checkfalse |
| Manual- plasma and red cell reduction  | checkfalse |
| RBC/plasma reduction with HES  | checkfalse |
| Ficoll sedimentation   | checkfalse |
| Centrifugation and volume reduction  | checkfalse |
| No processing  | checkfalse |
| Manual- plasma reduction only  | checkfalse |
| Other,   | checkfalse |
| The current pre-freeze processing method(s):   |            |
| AXP  | checkfalse |
| SEPAX  | checkfalse |
| Optipress  | checkfalse |
| Prepacyte  | checkfalse |
| Manual- plasma and red cell reduction  | checkfalse |
| RBC/plasma reduction with HES  | checkfalse |
| Ficoll sedimentation   | checkfalse |
| Centrifugation and volume reduction  | checkfalse |
| No processing  | checkfalse |
| Manual- plasma reduction only  | checkfalse |
| Other,   | checkfalse |
| Additives currently in use in addition to anticoagulants and DMSO during processing: |            |
| HES  | checkfalse |
| Prepacyte  | checkfalse |
| Plasmalyte   | checkfalse |
| Albumin  | checkfalse |
| Isotonic salt solution NaCl (saline)   | checkfalse |
| No additive  | checkfalse |
| Other,   | checkfalse |
| Cryoprotectant additives currently in use:   |            |
| Cryopreservation method currently in use:  |            |
| BioArchive   | checkfalse |
| MVE 1850 Vapor freezer   | checkfalse |
| Programmed cryopreservation with Air Liquid program (FREEZAL)                        | checkfalse |
| Programmed freezer   | checkfalse |
| Two-step (equilibrium) freezing  | checkfalse |

|  |            |
|--|------------|
| Cryobag currently in use:  |            |
| Currently used packaging when a unit is stored:                  |            |
| Segments currently stored with the unit by the cord blood bank:  |            |
| One attached segment   | checkfalse |
| Two or more attached segments                                    | checkfalse |
| Separate segments detached from the bag, but stored with the CBU | checkfalse |
| Other samples  | checkfalse |
| None   | checkfalse |

## Section 7 Testing

| Question  | CBB answer |
|---|------------|
| Does your cord blood bank currently store extra material? |            |
| Cord blood DNA  | checkfalse |
| Cord blood material for DNA extraction                    | checkfalse |
| Plasma/cord blood   | checkfalse |
| Maternal DNA  | checkfalse |
| Maternal material for DNA extraction                      | checkfalse |
| Maternal plasma/serum                                     | checkfalse |
| HLA typing at time of listing:                            |            |
| HLA-A   |            |
| HLA-B   |            |
| HLA-C   |            |
| HLA-DRB1  |            |
| HLA-DRB2  |            |
| HLA-DPB1  |            |

## Section 8 Storage

| Question   | CBB answer |
|--|------------|
| The following type(s) of storage container is currently used by the cord blood bank: |            |
| BioArchive tank  | checkfalse |
| Conventional storage tank-Vapor phase  | checkfalse |
| Conventional tank-Liquid phase   | checkfalse |
| Double walled liquid Nitrogen  | checkfalse |
| Type following type(s) of storage monitoring is currently by the cord blood bank:    |            |
| Alarm on individual tanks only   | checkfalse |
| Centralized system-local   | checkfalse |
| Centralized system-remote monitoring   | checkfalse |

|                           |            |
|---------------------------|------------|
| LN2 level                 | checkfalse |
| Lid opening               | checkfalse |
| System default            | checkfalse |
| Temperature monitoring    | checkfalse |
| No temperature monitoring | checkfalse |

## Section 9 Adverse Events Reporting

| Question   | CBB answer |
|--|------------|
| Adverse Event Reporting used by the cord blood bank: |            |
| Competent authority                                  | checkfalse |
| Internal report                                      | checkfalse |
| National registry                                    | checkfalse |
| Transplant centre                                    | checkfalse |
| WMDA   | checkfalse |