



## Donor Collection Data

### DONOR DATA

for **related** donors here:

Related-ID:

for **unrelated** donors here:

GRID:

### DONOR CONSENT

Donor agrees that her/his encoded data will be transmitted to the database of the EBMT (European Group for Blood and Marrow Transplantation).

She/he has signed the informed consent: ☐ yes ☐ no Date:

### DONOR DATA

Donor date of birth:

Sex: ☐ male ☐ female

Initials (*first name, last name*):

Language: please select

☐ **unrelated** donor

☐ **related** donor → relationship to recipient:

☐ Syngeneic (identical twin)

☐ Sibling

☐ Haplo, if yes specify: please select

☐ Other family member, specify:

### RECIPIENT DATA

UPN (only for **related** patient):

Patient ID (only for **unrelated** patient):

Initials (*first name, last name*):

Recipient date of birth:

Country of transplant:

### REPORT DATA

Date of this report:

Collection Center:

### PRODUCT

☐ Bone Marrow (BM) ☐ Mesenchymal Stem Cells (MSC)

☐ Peripheral Blood Stem Cells (PBSC) ☐ Unstimulated leucapheresis for DLI

☐ Unstimulated leucapheresis for virus-specific T-cells ☐ Whole blood donation for DLI



<b>related:</b> Related-ID: UPN:	<b>unrelated:</b> GRID: Patient ID:																						
<b>DONOR EVALUATION BEFORE DONATION</b> Date of evaluation: _____ Is a coexisting disease or organ impairment present at the time of evaluation/donation? <input type="checkbox"/> yes <input type="checkbox"/> no if yes, please specify: <table border="0"><tr><td><input type="checkbox"/> Cardiovascular</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Pulmonary</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Gastrointestinal</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Genito-urinary</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Neurological</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Immune/autoimmune</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Infectious</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Haematological</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Oncological</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Psychological</td><td>ICD 10 Code:</td></tr><tr><td><input type="checkbox"/> Other, please specify:</td><td>ICD 10 Code:</td></tr></table>		<input type="checkbox"/> Cardiovascular	ICD 10 Code:	<input type="checkbox"/> Pulmonary	ICD 10 Code:	<input type="checkbox"/> Gastrointestinal	ICD 10 Code:	<input type="checkbox"/> Genito-urinary	ICD 10 Code:	<input type="checkbox"/> Neurological	ICD 10 Code:	<input type="checkbox"/> Immune/autoimmune	ICD 10 Code:	<input type="checkbox"/> Infectious	ICD 10 Code:	<input type="checkbox"/> Haematological	ICD 10 Code:	<input type="checkbox"/> Oncological	ICD 10 Code:	<input type="checkbox"/> Psychological	ICD 10 Code:	<input type="checkbox"/> Other, please specify:	ICD 10 Code:
<input type="checkbox"/> Cardiovascular	ICD 10 Code:																						
<input type="checkbox"/> Pulmonary	ICD 10 Code:																						
<input type="checkbox"/> Gastrointestinal	ICD 10 Code:																						
<input type="checkbox"/> Genito-urinary	ICD 10 Code:																						
<input type="checkbox"/> Neurological	ICD 10 Code:																						
<input type="checkbox"/> Immune/autoimmune	ICD 10 Code:																						
<input type="checkbox"/> Infectious	ICD 10 Code:																						
<input type="checkbox"/> Haematological	ICD 10 Code:																						
<input type="checkbox"/> Oncological	ICD 10 Code:																						
<input type="checkbox"/> Psychological	ICD 10 Code:																						
<input type="checkbox"/> Other, please specify:	ICD 10 Code:																						
<b>DONATION PROCEDURE</b> Even if the preparative actions (i.e. start of injections, apheresis or anesthesia) are stopped prematurely (due to donor or recipient reasons) the activity fulfills the definition of a donation procedure and the donor should be registered and followed. PBSC: Date of 1 <sup>st</sup> injection (mobilisation): _____ BM: Start of anaesthesia: _____ Unstimulated leucapheresis (e.g. DLI): Start of apheresis: _____ Chronological number of this donation procedure: _____ If >1: for same recipient: <input type="checkbox"/> yes <input type="checkbox"/> no Center of previous donation: _____ Date of previous donation: _____																							
<b>DONATION DATA (1/2)</b> Date of first day of this collection: _____ Was the product collection completed? <input type="checkbox"/> yes <input type="checkbox"/> no If no, please specify: _____ Were hematopoietic growth factors used (e.g. G-CSF)? <input type="checkbox"/> yes <input type="checkbox"/> no If yes, please specify brand name: <input type="checkbox"/> Neupogen <input type="checkbox"/> Granocyte <input type="checkbox"/> Neulasta <input type="checkbox"/> Filgrastim Teva <input type="checkbox"/> Zarzio <input type="checkbox"/> other: _____																							



<b>related:</b> Related-ID: UPN:	<b>unrelated:</b> GRID: Patient ID:
<b>DONATION DATA (2/2)</b> Total dose per injection: _____ µg/kg (please convert I.E. to µg) Number of doses per day: _____ Total number of doses: _____ Were cell binding inhibitors (e.g. Plerixafor) used? <input type="checkbox"/> yes <input type="checkbox"/> no Was erythropoietin used? <input type="checkbox"/> yes <input type="checkbox"/> no Were other drugs used for mobilisation? <input type="checkbox"/> yes <input type="checkbox"/> no if yes, please specify: Donor weight: _____ kg	
<b>APHERESIS COLLECTION</b> Number of apheresis performed: _____ Collection technique: <input type="checkbox"/> by peripheral veins <input type="checkbox"/> by central venous catheter	
<b>BONE MARROW COLLECTION</b> Anaesthesia: <input type="checkbox"/> general <input type="checkbox"/> epidural/spinal <input type="checkbox"/> local anaesthesia Autologous blood donation prior to collection? <input type="checkbox"/> yes <input type="checkbox"/> no Was autologous blood re-transfused? <input type="checkbox"/> yes <input type="checkbox"/> no	
<b>COMPLICATIONS</b> Report every Serious Adverse Reaction or Serious Adverse Event (SARE) occurring within the interval between start and end of the donation procedure with ICD coding. Serious Adverse Reaction or Serious Adverse Event (SARE): <input type="checkbox"/> yes <input type="checkbox"/> no if yes, please specify: ▪ ICD 10 Code: _____ Date: _____ ▪ ICD 10 Code: _____ Date: _____ Comments:  Report any Adverse Event or Adverse Reaction occurring within the interval between start and end of the donation procedure, which is noteworthy and serves as important information for FollowUp SBSC. Adverse Reaction or Adverse Event: ▪ In treatment: <input type="checkbox"/> Treatment completed: <input type="checkbox"/> ▪ In treatment: <input type="checkbox"/> Treatment completed: <input type="checkbox"/> Comments:	
<b>DONOR CARE</b> <input type="checkbox"/> as out-patient <input type="checkbox"/> as in-patient	

Form completed by (first/last name):

Date: