| Donor last name | Donor first name | Donor ID | an_gridformatted |
|-----------------|------------------|----------|------------------|
| lastname | firstname | | |
| | | | |

CONSENT TO DONATE STEM CELLS FROM A BONE MARROW COLLECTION

The original consent form should be retained by the Collection Centre. One copy should then be retained by the donor and a copy forwarded to Anthony Nolan.

A. STATEMENT BY HEALTHCARE PROFESSIONAL (Please tick the boxes)

I confirm that the donor for whom consent is being taken has identified themselves by confirming their name, date of birth and home address information supplied to me by Anthony Nolan.

I have explained the proposed procedure of a bone marrow stem cell collection to the volunteer donor and briefly discussed the intended benefits to the patient. In particular, I have explained to the donor:

- 1. the need for microbiology and virology testing and in particular the need to test the donor's blood for markers of infection including syphilis, HIV, HTLV, and Hepatitis B, C & E
- 2. the need for a general anaesthetic and any possible serious or frequently occurring side effects from this procedure
- **3.** the process of bone marrow aspiration and any serious or frequently occurring risks or side effects that may be involved in the procedure
- **4.** the need to be admitted to hospital for two nights and to rest at home for up to 2 weeks after discharge to help recovery and to reduce, where possible, side effects following the procedure
- 5. the possible short and long-term risks associated with donating bone marrow stem cells including:
 - after the procedure it is expected for haemoglobin to be lower, because the bone marrow contains many red cells. In some cases, they may become anaemic. I have explained that there may be a need of oral iron or if the pre-harvest ferritin is low, IV iron may be necessary on the day of admission for the bone marrow harvest
 - after the procedure it is normal to have some pain at the aspiration site which is usually well-controlled with oral pain killers (analgesia). Approximately 5-10 in 100 donors **may** have ongoing pain lasting up to 4 weeks. I have explained that long-term pain has been reported in approximately 1-2 in 100 donors.
 - surgical wounds will be present after the procedure (between 1 3 puncture sites) on each side of the lumbar vertebrae)
 - the major risk of donating bone marrow stem cells is associated with anaesthesia and includes (uncommon & extremely rare) the following: aspiration pneumonia, pulmonary embolus, arrythmias, cerebral infarction, allergy and cardiac arrest. The risk of death due to a general anaesthetic is <1/10,000.
 - the specific procedure related risks: bleeding, low blood pressure, bacteraemia, local infection and/or haematomas (bruises) at the harvest puncture sites, post-operative fever, fractured iliac crests, and in extremely rare cases: broken aspiration needles requiring surgical removal, transient pressure neuropathies (numbness) spinal headache and bone marrow or air emboli.
 - the possibility (low risk) that a blood transfusion may be required during or after the procedure.
- **6.** To reduce risk of possible exposure to transmissible infections ahead of donation, including unprotected sex with a new or high-risk sexual partner or intravenous drug use, and if such activity occurs to inform Anthony Nolan to facilitate further testing
- 7. the requirement to store confidential information in accordance with applicable data protection and related laws and guidance (see section F below)

| Don | or last name | Donor first name | Donor ID | an_gridformatted | |
|--|--|----------------------------|---------------------|------------------|--|
| lastr | name | firstname | | | |
| | | | | | |
| 8. the possible storage of cells, the need for discard of stored material as well as the possible use of cells for research purposes by the transplant centre or Anthony Nolan (which depending on the circumstances, may be outside of the UK and the EEA) ("the Transplant Centre"). | | | | | |
| 9. | that a copy of all test result | s and findings will be ser | nt to Anthony Nolar | า | |
| 10. | 10. the potential need for cryopreservation should the transplant centre request this for patient safety | | | | |
| Pleas | lease tick this box to confirm you have explained points 1 to 10 above to the donor | | | | |
| | Please tick this box to confirm you believe the donor understands the information provided and can freely give consent | | | | |
| The current versions of the HTA's Codes of Practice on the Donation of Allogeneic Bone Marrow and Peripheral Blood Stem Cells for Transplantation, and on Consent The current version of the HTA's Guidance for Transplant Teams and Accredited Assessors and have applied the principles and procedures accordingly. | | | | | |
| Signe | ed by Healthcare Profession | nal | Date of assessmer | nt | |
| First | name | | Last name | | |
| Job t | itle | | Collection centre | | |

| Donor last name lastname | Donor first name firstname | Donor ID an_gridformatted | |
|---|---|--|-------------------------------|
| B. STATEMENT BY DONOR: PRO | OCEDURE INFORMATION (| Please tick the boxes) | |
| I've been told I'm a match for a pa asked to donate haematopoietic chosen to donate my cells throug collection | (blood) stem cells. After cons | sideration I've voluntarily | |
| The healthcare professional name | ed in section A has clearly exp | lained to me: | |
| • the donation procedure, in | ncluding the general anaesth | etic | |
| • the possible short and lon | g-term risks related to the co | llection | |
| | ke extra precautions ahead o n that could be passed to the | f my donation to reduce the risk patient | |
| if I have any new sexual pa Nolan via my coordinator | rtners between now and the o | donation, to inform Anthony | |
| I have received and understood the opportunity to ask questions. Any obeen given sufficient information to | questions have been answere | ed to my satisfaction. I believe I hav | 'e |
| evidence of important infe viruses. I understand that it | ections including those cause f the results of any of these te | and to check that my blood does noted by the syphilis, HIV, HTLV, and he ests are abnormal, I will be informed bllow-up will be arranged by Anthor | patitis B, C & E d. I also |
| undergo a general anaesth marrow transplant | netic for the purpose of dona | ting marrow for a patient requiring | a bone |
| 3. donate the necessary amo | ount of my bone marrow to a p | patient | |
| Please tick this box to confirm you | ur agreement with points 1 to | 3 above | |
| lunderstand that: | | | |
| | re to discuss and consider thi | nis patient on a second occasion. I a s, but also understand that I am fre | |
| the donor collection centr | e. The basic risks to the patie ening implications for the pat | th my Anthony Nolan coordinator on the have been explained to me and I lient if I withdraw after the patient h | fully |
| Please tick this box to confirm you | ur agreement with points 4 to | 5 above | П |

| Donor last name lastname | Donor first name firstname | Donor ID | an_gridformatted |
|--|--|-----------------|--|
| n addition, l understand that: | | | |
| 6. I cannot be given a guaran procedure, although the h | tee that a specifically named ealthcare professional will ha | • | • |
| 7. I will be given further oppoprocedure | ortunity to discuss the details o | of anaesthes | ia with an anaesthetist before the |
| | | | cipate in routine follow-ups post- t eight and 10 years after donation. |
| 9. my stem cells will be given who may remain anonymo | | y will be main | tained for at least two years, and |
| 10. the patient who receives n | ny cells may be of any age, rac | ce or religion | and be living in any part of the world |
| 11. the primary responsibility staff who undertake the p | | on rests with t | he medical and other professional |
| 12. this consent is automatica collection | ally cancelled if I am found not | to be fit to d | onate stem cells by bone marrow |
| 13 Transplant is carried out in | the hope that it will cure the p | patient Sadly | however the patient may not be |

cured and may not survive in the longer-term

Please tick this box to confirm your agreement with points **7** to **13** above

| Donor last name lastname | Donor first name firstname | Donor ID | an_gridformatted |
|-----------------------------|----------------------------|----------|------------------|
| | | | |

C.STATEMENT BY DONOR: STORAGE, USE AND DISCARD OF CELLS AT TRANSPLANT CENTRE

I understand that:

- 1. some of my blood, cells or DNA (which may be taken from blood or cells provided by me prior to, or at the time of, donation) may be stored for the purposes of undertaking tests to monitor and appropriately treat the patient of this particular transplant
- 2. a small part of my donation may be stored as a source of therapeutic cells to be administered to the patient after the transplant if needed
- **3.** fresh or frozen samples of my blood, cells or DNA may be used for the purposes of quality control monitoring, clinical audit, public health surveillance purposes and/or future testing relevant to the quality of my stored cells
- **4.** my cells will be disposed of, when they are no longer required or prove unsuitable for clinical use (or for research, if I have provided consent), in a manner which meets applicable regulations for the disposal of biohazardous materials

| Please tick this box to confirm your agreement with points 1 to 4 above | П |
|---|-----|
| ricase der diis box to commini your agreement with points 1 to 4 above | 1 1 |

Further testing of the cells following infusion to the patient

lunderstand that:

please tick here

- Following my cells being infused into the patient, the transplant centre may carry out testing to support the patient's recovery. These tests may include genetic screening, as well as screening for other blood disorders. These tests are performed on the patient and not directly on my blood samples. However, as the patient's blood cells are made from donor stem cells, in rare cases these tests may find a genetic variant thought to have originated from donor cells, your cells.
- Confirming if a genetic variant originated from donor stem cells or recipient cells is not possible without additional confirmatory genetic testing.
- Some genetic findings of potential donor origin may be relevant to my health and wellbeing, or the health and well-being of my children (or future children).
 - 1. any genetic findings from the patient, thought to have originated from donor cells, that are considered of clinical significance, or are of uncertain significance at the time of testing, will be shared with me in order to arrange appropriate genetic counselling and testing, which I can accept or decline.
 - 2. any genetic findings from the patient, thought to have originated from donor cells, that are not considered of clinical significance, or are of uncertain significance at the time of testing, will not be routinely shared with me.

| Please tick this box to confirm your agreement to being informed of any clinically significant findings of potential donor origin (points ${f 1}$ and ${f 2}$) | П |
|---|---|
| OR | |
| If you do not want to be informed of any clinically significant genetic findings of potential donor origin, even if life-threatening or preventable conditions or when withholding information may be harmful, | |

П

| lastname firstname | |
|--------------------|--|
| | |
| | |

D. STATEMENT OF DONOR: CRYOPRESERVATION OF BONE MARROW DONATION

On occasion, a transplant centre may request to freeze (cryopreserve) the donated stem cells, to be infused to the patient on a later date. This may be due to patient issues, scheduling or logistics issues.

In addition to consenting to the donation procedure in the terms set above in section B:

- 1. I voluntarily consent to the cryopreservation of my cells, if necessary, and understand that the stem cells collected during this bone marrow donation process may be cryopreserved for infusion at a later date
- 2. If my cells are cryopreserved, I give consent for my cells to be discarded if they are no longer required or prove unsuitable for clinical or research use, and in this event, I will be informed by Anthony Nolan
- **3.** If discarded, I understand they will be disposed of appropriately according to applicable regulations for the disposal of biohazardous materials

| Please tick this box to confirm your agreement with points 1 to 3 above | |
|---|--|
| OR | |
| I do not consent to my cells being cryopreserved | |

| firstname | | |
|-----------|----------|----------|
| | | |
| fi | irstname | irstname |

E. STATEMENT BY DONOR: USE OF CELLS FOR RESEARCH

On occasion, there may be cells remaining in the product bag post-transplant and Anthony Nolan or transplant centres may request to use these remaining cells for research purposes. This may also be the case with the full donation if, for any reason, the transplant cannot take place. In these cases, requests are assessed and approved by a properly constituted research ethics committee and undertaken in accordance with appropriate ethical, legal and professional standards.

lunderstand that:

- 1. Some or all of my blood, cells or DNA from this collection could be used in a non-identifiable way for future medical research projects. I will not benefit financially from any research undertaken and I waive all rights to any registered patents
- 2. My participation in the storage of my blood, cells or DNA for research is voluntary. Refusal to participate will not affect my status on the Anthony Nolan register as a stem cell donor or result in the loss of any benefits such as follow-up care following my donation
- **3.** My pseudonymised data may be used to support such research and will be used in accordance with the Anthony Nolan Privacy Policy
- 4. I have the right to withdraw consent for the use of my blood, cells or DNA for research without it affecting my status on the Anthony Nolan register as a stem cell donor or resulting in the loss of any benefits, such as follow-up care post-donation. I understand that once my cells have been used for a research study, they will not be able to be withdrawn from that study.

| they will het be able to be with a awill of the third to ady. | |
|--|--|
| Please tick this box to confirm your agreement with points 1 to 4 above | |
| OR | |
| Please tick this box to confirm that you do not want your blood, cells or DNA to be used for future research | |

| Donor last name | Donor first name | Donor ID | an_gridformatted |
|-----------------|------------------|----------|------------------|
| lastname | firstname | | |
| | | | |

F. STATEMENT BY DONOR: PRIVACY

| give my consent to Anthony Nolan processing and storing the following data as per the Anthony No policy (available at anthonynolan.org/privacy), specifically: | lan privac |
|---|------------|
| The data I have provided in this form | |
| Any analysis of the blood samples I provide, which I understand will be tested for markers of infection including syphilis, HIV, HTLV and Hepatitis B, C $\&$ E | |
| The results of blood tests, which I specifically consent to Anthony Nolan sharing with my GP, if deemed necessary for medical reasons | |
| Any analysis of the stem cells I donate, which I understand may be stored by the transplant centre and/or Anthony Nolan for patient transplant and, if I have agreed, for research purposes | |
| All health and medical information I provide, which I understand may be stored by the transplant centre and Anthony Nolan in order to establish I am medically fit to donate for a patient | |
| My pseudonymised personal data that may be shared with third party organisations including but not limited to the European Group for Blood and Marrow Transplant registry, to analyse factors that contribute to the outcome of transplants, in accordance with applicable data protection and related laws and guidance | |
| I understand that if the patient is based outside of the UK, my personal data will be shared with an international donor registry and/or international transplant centre in accordance with the Anthony Nolan Privacy Policy | |
| I consent to Anthony Nolan's transfer of my data (in pseudonymised form) to countries without the same data protection laws as the UK/EU for the purposes stated in the Anthony Nolan privacy policy. Anthony Nolan agrees to protect my data as described in its Privacy Policy and provide adequate protection for transfers to countries outside the UK and EEA. | |
| I understand that I have the right to access my medical information in accordance with applicable data protection and related laws and guidance | |

| Donor last name | Donor first name | Donor ID | an_gridformatted |
|-----------------|------------------|----------|------------------|
| lastname | firstname | | |
| | | | |

| DONOR I confirm that I have read and completed parts B, C, D, E and F of this form. | | | | |
|---|--|--|--|--|
| Signed by Donor | Date | | | |
| Donor first name | Donor last name | | | |
| HEALTHCARE PROFESSIONAL I confirm that I have witne of this form. | ssed the above donor completing parts B, C, D, E and F | | | |
| Signed by Healthcare Professional (usually same individual in section A) | Date | | | |
| Healthcare Professional first name | Healthcare Professional last name | | | |
| Healthcare Professional title (and email if not the Healthcare Pr | ofessional mentioned in section A) | | | |

| Donor last name lastname | Donor first name firstname | Donor ID | an_gridformatted |
|-----------------------------|----------------------------|----------|------------------|
| | | | |

G. CONFIRMATION OF CONSENT

| TO BE COMPLETED BY THE DONOR AND HEALTHCARE PROFESSIONAL WHEN THE DONOR IS ADMITTED FOR THE PROCEDURE | | | | |
|--|-----------------------------------|--|--|--|
| DONOR please tick the relevant box | | | | |
| I confirm that I have no further questions and that I wish to donation. I confirm that I have not been coerced, paid, or received this donation. | | | | |
| OR | | | | |
| I withdraw my consent and will not be proceeding | | | | |
| Signed by Donor | Date | | | |
| Donor first name | Donor last name | | | |
| Healthcare Professional | | | | |
| Signed by Healthcare Professional | Date | | | |
| Healthcare Professional first name | Healthcare Professional last name | | | |
| Job title | Collection centre | | | |