Donor last name lastname	Donor first name Firstname	Donor ID an_gridformatted

CONSENT TO DONATE STEM CELLS FROM A BONE MARROW COLLECTION (RELATED)

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)

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 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 results and findings will be sent to Anthony Nolan 10. the potential need for cryopreservation should the transplant centre request this for patient safety 			
Please tick this box to confirm you	u have explained points 1 to 10	above to the donor	
Please tick this box to confirm you and can freely give consent	lease tick this box to confirm you believe the donor understands the information provided and can freely give consent		
	erstood: e HTA's Codes of Practice on a and Peripheral Blood Stem Ce		
The current version of the HTA's Guidance for Transplant Teams and Accredited Assessors and have applied the principles and procedures accordingly.			
Signed by Healthcare Profession	nal Date	of assessment	
First name	Last n	ame	
Job title	Collec	ction centre	

Donor ID

Donor first name

Donor last name

lastn	ame	Firstname	an_gridformatted	
D CTA	TEMENT BY DONOR, DD	OCEDURE INFORMATION (Plea		
		·	•	
asked	to donate haematopoietic n to donate my cells throug	itient in need of a bone marrow tra (blood) stem cells. After consider In the procedure known as a bone	ation I've voluntarily	
The he	ealthcare professional name	ed in section A has clearly explaine	d to me:	
•	the donation procedure, i	ncluding the general anaesthetic		_
•	the possible short and lon	g-term risks related to the collect	ion	
•	•	ke extra precautions ahead of my n that could be passed to the patio		
•	if I have any new sexual pa Nolan via my coordinator	rtners between now and the dona	tion, to inform Anthony	
opport	unity to ask questions. Any	e information provided to me by A questions have been answered to o give my informed consent to pro	my satisfaction. I believe I ha	ve
1.	evidence of important infeviruses. I understand that i	certain my fitness to donate and to ections including those caused by f the results of any of these tests a sts, counselling, and clinical follow	the syphilis, HIV, HTLV, and he re abnormal, I will be informe	epatitis B, C & E d. I also
2.	undergo a general anaesti marrow transplant	netic for the purpose of donating r	narrow for a patient requiring	j a bone
3.	donate the necessary amo	ount of my bone marrow to a patie	nt	
Please	e tick this box to confirm you	ur agreement with points 1 to 3 abo	ove	
lunders	stand that:			
4.	. ,	be asked to donate cells to this pa re to discuss and consider this, bu tion at any time		•
5.	the donor collection centr	ets at any time by speaking with my re. The basic risks to the patient ha ening implications for the patient i ent conditioning treatment	ve been explained to me and	Ifully
Please	e tick this box to confirm you	ur agreement with points 4 to 5 ab	ove	

Donor first name

Donor ID

Donor last name

Donor last name	Donor first name	Donor ID
lastname	Firstname	an_gridformatted

In addition, I understand that:

- **6.** I cannot be given a guarantee that a specifically named healthcare professional will perform the procedure, although the healthcare professional will have the required training and experience
- 7. I will be given further opportunity to discuss the details of anaesthesia with an anaesthetist before the procedure
- **8.** my recovery will be monitored by Anthony Nolan and I agree to participate in routine follow-ups post-donation, as well as annually up to six years. Follow-ups will then be at eight and 10 years after donation.
- **9.** the primary responsibility for the bone marrow collection rests with the medical and other professional staff who undertake the procedure
- 10. this consent is automatically cancelled if I am found not to be fit to donate stem cells by bone marrow collection
- 11. Transplant is carried out in the hope that it will cure the 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 11 above	
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Donor last name lastname	Donor first name Firstname	Donor ID an_gridformatted
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C.STATEMENT BY DONOR: STORAGE, USE AND DISCARD OF CELLS AT TRANSPLANT CENTRE

lunderstand 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	
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Further testing of the cells following infusion to the patient

lunderstand that:

- 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 ${f 2}$)	
C	DR .	_
	If you <u>do not</u> 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, please tick here	

Donor last name	Donor first name	Donor ID
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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 ${f 1}$ to ${f 3}$ above	
OR	
I <u>do not</u> consent to my cells being cryopreserved	П

Donor last name	Donor first name	Donor ID
lastname	Firstname	an_gridformatted

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.

Please tick this box to confirm your agreement with points 1 to 4 above	
OR	
Please tick this box to confirm that you <u>do not</u> want your blood, cells or DNA to be used for future research	

Donor last name	Donor first name	Donor ID
lastname	Firstname	an_gridformatted

F. STATEMENT BY DONOR: PRIVACY

give my consent to Anthony Nolan processing and storing the following data as per the Anthongoolicy (available at anthonynolan.org/privacy), specifically:	y Nolan privacy
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	
I understand that if clinically relevant for the patients' health, my health and medical information may be shared between the transplant centre and 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
lastname	Firstname	an_gridformatted

G. DONOR AND HEALTHCARE PROFESSIONAL DECLARATION

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

 $HEALTHCARE\ PROFESSIONAL: I confirm\ that\ I have\ witnessed\ the\ above\ donor\ completing\ parts\ B,\ C,\ D,\ E\ and\ F\ of\ this\ form.$

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 Professional mentioned in section A)		

Donor last name	Donor first name	Donor ID		
lastname	Firstname	an_gridformatted		
H. 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 proceed with stem cell donation. I confirm that I have not been coerced, paid, or received any inducement in relation to П 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