Donor last name	Donor first name	DonorID	an_gridformatted
lastname	firstname		

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

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 the procedure, 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)

lastname	firstname	Donorid an_gridiormatted		
 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 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 believe the donor understands the information provided and can freely give consent]	
I confirm that I have read and unde	erstood:			
	e HTA's Codes of Practice o and Peripheral Blood Stem (
 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. 				
Signed by Healthcare Professional Date of assessment				
First name	Las	t name		
Job title	Col	Collection centre		

Dono	or last name ame	Donor first name firstname	DonorID	an_gridformatted	
B. STA	TEMENT BY DONOR: PRO	OCEDURE INFORMATION (Please tick t	he boxes)	
asked	to donate haematopoietic n to donate my cells throug	atient in need of a bone marro (blood) stem cells. After con gh the procedure known as a b	sideration I'v	e voluntarily	
The he	ealthcare professional name	ed in section A has clearly exp	lained to me:		
•	the donation procedure, i	ncluding the general anaesth	etic		
•	the possible short and lor	ng-term risks related to the co	ollection		
•	-	ake extra precautions ahead c on that could be passed to the	-	n to reduce the risk	
•	if I have any new sexual pa Nolan via my coordinator	artners between now and the	donation, to i	nform Anthony	
opport	unity to ask questions. Any	e information provided to me questions have been answere to give my informed consent t	ed to my satis	faction. I believe I ha	ve
1.	evidence of important info viruses. I understand that i	certain my fitness to donate a ections including those cause f the results of any of these te sts, counselling and clinical fo	ed by the syplests are abno	nilis, HIV, HTLV, and he rmal, I will be informe	epatitis B, C & E d. I also
2.	undergo a general anaest marrow transplant	hetic for the purpose of dona	ting marrow t	for a patient requiring	ga bone
3.	donate the necessary amo	ount of my bone marrow to a p	oatient		
Please	e tick this box to confirm yo	ur agreement with points 1 to	3 above		
lunders	stand that:				
4.		be asked to donate cells to the reto discuss and consider thing at any time	•		•
5.	the donor collection centrunderstand the life-threat	nts at any time by speaking with re. The basic risks to the patient rening implications for the pata ant conditioning treatment	nt have been	explained to me and	Ifully
Please	e tick this box to confirm yo	ur agreement with points 4 to	5 above		
In addit	ion, l understand that:				

Dono lastna	or last name ame	Donor first name firstname	DonorID	an_gridformatted
6.		tee that a specifically named ealthcare professional will ha	•	•
7.	I will be given further oppo procedure	ortunity to discuss the details	of anaesthes	sia with an anaesthetist before the
8.			•	cipate in routine follow-ups post- at eight and 10 years after donation.
9.	my stem cells will be given who may remain anonymo	•	y will be mair	ntained for at least two years, and
10.	. the patient who receives r	ny cells may be of any age, ra	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	the medical and other professional
12.	this consent is automatica collection	ally cancelled if I am found not	to be fit to d	lonate stem cells by bone marrow
13.	transplant is carried out in cured and may not survive	•	atient. Sadly	however, the patient may not be

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

Donor last name	Donor first name	Donor ID	an_gridformatted
lastname	firstname		

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:

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- 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.

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 1 and 2)	
R	
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	DonorID	an_gridformatted
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
- If discarded, Lunderstand they will be disposed of appropriately according to applicable regulations for

•	the disposal of biohazardous materials	uons i
	Please tick this box to confirm your agreement with points 1 to 3 above OR	
	I do not consent to my cells being cryopreserved	

Donor last name	Donor first name	Donor ID	an_gridformatted
lastname	firstname		

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	an_gridformatted
lastname	firstname		

F. STATEMENT BY DONOR: ANTHONY NOLAN PATIENT DONOR PROJECT

Anthony Nolan is undertaking a research study that we would like you to consider joining. This study is investigating the importance of HLA matching (tissue typing) and other genetic factors that have been shown to influence the outcome of unrelated stem cell transplants.

Although this research will not directly alter results in this specific transplant, it is hoped that in future it will enable us to advise which donor should be chosen in the event that no fully matched donor is available, but where there is a choice of partially matched donors.

We are asking UK donors and all patients who receive stem cells from a UK donor to join this research project. The DNA extracted from this sample will only be used for matching studies in our laboratory (i.e. only looking for factors to do with outcome in haematopoietic stem cell transplants). It will be stored within the Research Institute, with a unique coding number for the duration of the study (i.e. only the researchers will be able to link the sample to the person who provided it).

After the study is completed, we would like to store the donor/patient sample pairs in a anonymised form (i.e. the details cannot be traced back to an individual person). The purpose of this is to enable us to test these samples for any genetic factors related to stem cell transplantation that may be discovered in years to come. These samples will be owned by Anthony Nolan. All that will be required from you will be a blood and/or a buccal swab sample (mouth swab). If you choose not to join this study, it will not affect your treatment/donation in any way.

lunderstand the following:

- 1. I have read and fully understood the above information regarding participating in an Anthony Nolan research study.
- 2. I have had the opportunity to ask questions and have received satisfactory answers.
- **3.** my participation is voluntary and if I choose not to provide a blood and/or buccal cell sample (mouth swab), my treatment/donation will not be affected in any way.
- 4. Lagree to take part in the study by providing a blood and/or buccal cell sample (mouth swab).
- **5.** I agree that my blood and/or buccal cell sample (mouth swab) can be retained after the study completes (in a anonymised form).
- **6.** Anthony Nolan will use and store my personal data in accordance with the Anthony Nolan Privacy Policy and that I may withdraw my consent to the use of my personal data, at any time, in accordance with the terms of this policy.

Please tick this box to confirm your agreement with points 1 to 6 above	
OR	
I do not want to be part of this study	

Donor last name	Donor first name	DonorID	an_gridformatted
lastname	firstname		

G. STATEMENT BY DONOR: PRIVACY

I 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 p	orivacy
The data I have provided in this form		
Any analysis of the blood sample I donate, which I understand will be tested for markers of infection including syphilis, HIV, HTLV, and Hepatitis B, C $\&$ E		
The results of such 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 to 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 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	. 🗆	
I understand that I have the right to access my medical information in accordance with applicable data protection and related laws and guidance		
Additional statement only relevant to participants in the Anthony Nolan Patient Donor Project:		
Additionally, and only where I have agreed to participate in the research detailed in Section F, I give my consent to Anthony Nolan to use the data provided in this form and a sample of my DNA for the purposes of the research outline at section F above.		

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 witned this form.	essed the above donor completing parts B, C, D, E and F
Signed by Healthcare Professional (usually same individual in section A)	Date
	Date Healthcare Professional last name

Donor last name	Donor first name	DonorID	an_gridformatted
lastname	firstname		

I. 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.	o proceed with stem cell	
I confirm that I have not been coerced, paid, or received any inducement in relation to this donation.		
OR		
I withdraw my consent and will <u>not</u> 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	