Donor last name	Donor first name	Donor ID	an_gridformatted
lastname	firstname		

CONSENT FORM FOR BLOOD STEM CELL DONATION

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 peripheral blood stem cell mobilisation and collection to the volunteer donor and briefly discussed the intended benefits to the patient. In particular, I have explained to the donor:

- 1. the use of Granulocyte-Colony Stimulating Factor (G-CSF) to mobilise the donor's stem cells and any serious or potential side effects from this drug
- 2. the need for microbiology testing and in particular the need to test the donor's blood for markers of infection including Syphilis, HIV, HTLV, Hepatitis B, C & E
- **3.** the use of a blood cell separator to collect the donor's stem cells and any serious or potential occurring side effects involved in the procedure
- 4. the potential need to insert a femoral, internal jugular or sub-clavian central venous line if peripheral access is not adequate, as well as any serious or frequently occurring risks associated with such a procedure. I have also explained that separate donor consent for this procedure would be required
- 5. the possible short and long-term risks associated with donating peripheral blood stem cells including:
 - hypocalcaemia (sudden drop of calcium in the bloods) due to the citrate (ACD-A) used in the
 apheresis procedure, which can cause transient paraesthesia (pins and needles, numbness),
 muscle spasms, cramps, and in severe untreated cases risk of seizures (extremely rare). This may
 require calcium tablets or occasionally IV calcium replacement
 - risks associated with G-CSF such as bone ache, myalgia, headache, fatigue, fever, chest pain and thrombocytopenia (low platelets). I have explained these will usually require analgesia(paracetamol)
 - that in extremely rare cases the following G-CSF side effects may occur; vascular event, splenic rupture, sore eyes and anaphylaxis (allergic reaction)
 - bruising and bleeding at the site of venepuncture or central line site
 - the possibility of infection of the venepuncture site
- **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)
- **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 (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

lastname	firstname				
10. the potential need for cryopreservation should the transplant centre request this for patient safety					
Please tick this box to confirm you h	ave explained points I to I	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 understood: 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. 					
Signed by Healthcare Professional Date of assessment		of assessment			
First name	Lastr	ame			
i ii striailie	Lasti	anie			
Job title Collection centre					

Donor ID an_gridformatted

Donor first name

Donor last name

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L						
ı	B. STATEMENT BY DONOR PROCEDURE INFORMATION (Please tick the boxes)					
	confir consideriple	een told I'm a match for a patie m compatibility, and I've been deration I've voluntarily choser neral blood stem cell collectio cells my body produces and th	nasked to donate haematop n to donate my cells through n (PBSC), which involves taki	oietic (blood the procedu ng a drug to	d) stem cells. After ire known as a mobilised	
	The he	ealthcare professional named the donation procedure, inc the administration of the dru	luding the use of a blood cel	l separator m	•	
	•	the possible short and long-	term risks related to the coll	ection		
	•	that if sexually active to take contracting an infection tha	•	•	to reduce the risk of	
	•	if I have any new sexual partr my coordinator	ners between now and the do	onation, to in	form Anthony Nolan via	
(I have received and understood the information provided to me by Anthony Nolan and have been given the opportunity to ask questions. Any questions have been answered to my satisfaction. I believe I have been given sufficient information to give my informed consent to proceed with the donation. I agree to:					
	1.	undergo blood tests to ascer evidence of important infecti viruses. I understand that if th understand that further tests, necessary	ons including those caused e results of any of these test	by the Syphil s are abnorm	lis, HIV, HTLV and Hepatit nal, I will be informed. I als	is B, C & E so
	2.	receive G-CSF in order to pro	oduce sufficient stem cells in	my circulatir	ng blood	
	3.	donate stem cells to a patien	t, collected by the use of the	apheresis m	nachine	
	Pleas	e tick this box to confirm your a	agreement with points 1 to 3	above		
	under	stand that:				
	4.	there is a possibility I may be a approached in the future to d request for a further donation	liscuss and consider this, but			
	5.	I may withdraw my consents a the donor collection centre. T understand the life-threateni commenced pre-transplant of	The basic risks to the patient ng implications for the patie	have been e	xplained to me and I fully	
	6.	following my cells being infus recovery. These tests may inc rare cases these tests may re- be contacted by Anthony Nol	clude genetic screening, as w sult in findings which may be	vell as screer	ning for other blood diso	rders. In
	Please	e tick this box to confirm your a	agreement with points 4 to 6	above		

Donor lastnar	last name me	Donor first name firstname	Donor ID	an_gridformatted
In additio	on, I understand that:			
		e that a specifically named he Ithcare professional will have		
				pate in routine follow-ups post- eight and 10 years after donation
	ny stem cells will be given to yho may remain anonymous		vill be mainta	ined for at least two years, and
10 . th	ne patient who receives my o	cells may be of any age, race	or religion ar	nd be living in any part of the world
	. , ,	the blood cell collection and hal staff who undertake the p		G-CSF therapy rests with the
	nis consent is automatically ell separator machine	cancelled if I am found not to	be fit to dor	nate blood stem cells using a blood
	ransplant is carried out in the ured and may not survive in 1	·	ient. Sadly h	owever, the patient may not be

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: STOR	AGE, USE AND DISCARD	OF CELLS AT TE	RANSPLANT CENTRE
lunderstand that:			
 some of my blood, cells or DN time of, donation) may be sto treat the patient of this partic 	red for the purposes of unde		
a small part of my donation m patient after the transplant if	•	herapeutic cells t	to be administered to the
fresh or frozen samples of my monitoring, clinical audit, pub of my stored cells			oses of quality control testing relevant to the quality
 my cells will be disposed of, v research, if I have provided co biohazardous materials 		•	uitable for clinical use (or for egulations for the disposal of
Please tick this box to confirm your a	agreement with points 1 to 4	above	
D. STATEMENT OF DONOR: CRY	OPRESERVATION OF PBS	SC DONATION	
On occasion, a transplant centre may the patient on a later date. This may b		· ·	
In addition to consenting to the dona	tion procedure in the terms s	set above in secti	on B:
I voluntarily consent to the cry collected during the PBSC do			
2. If my cells are cryopreserved, prove unsuitable for clinical o			
3. If discarded, I understand the the disposal of biohazardous		riately according	g to applicable regulations for

I do not consent to my cells being cryopreserved

OR

Please tick this box to confirm your agreement with points 1 to 3 above

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lastname firstn	name	

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 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

I give my consent to Anthony Nolan processing and storing the following data as per the Anthony N policy (available at anthonynolan.org/privacy), specifically:	olan 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	
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		

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			
$Health care\ Professional\ title\ (and\ email\ if\ not\ the\ Health care\ Professional\ mentioned\ in\ section\ A)$				

Donor last name lastname	Donor first name firstname	Donor ID an_gridformatted
H. CONFIRMATION OF CONSENT TO BE COMPLETED BY THE DONOR A ADMITTED FOR THE PROCEDURE		SIONAL WHEN THE DONOR IS
DONOR please tick the relevant box	(
I confirm that I have no further questi donation. I confirm that I have not been coerce this donation.		
OR		
I withdraw my consent and will not be	e proceeding	
Signed by Donor	Date	
Donor first name	Dono	r last name
Healthcare Professional	<u> </u>	
Signed by Healthcare Professional	Date	
Healthcare Professional first name	Health	ncare Professional last name
Job title	Collec	ction centre