

Loughborough University The Centre for Biological Engineering		Acquisition and Receipt of Biological Materials		
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ARF No: CBE/ARF/000 76

PART A: To be completed by the Receiver (a separate form must be completed for each sample type)

A1. Details of Sample/Specimen

Type/ID: (eg primary cell, cell line, tissue, body fluid, excreta, biological agent)	Human Cord Blood (Ficolled) CD34+ isolation	<input checked="" type="checkbox"/> Human <input type="checkbox"/> Animal
Format / Quantity: (eg vials, slides, etc)	3x50mL tube containing ~21mL ficol unit. 3 donors total	
Tissue site/Organ source:	Cord Blood	<input type="checkbox"/> N/A
Batch N°:		
Is the sample/specimen considered to be Relevant Material under the Human Tissue Act (HTA)? If No, go to section A2.	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	
Is the material obtained from an HTA licenced Tissue Bank with REC approval for generic research use?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	
Is the material obtained for storage and use under a project specific NHS REC approval?	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No	
Is the material licensable under the HTA? Indicate source below:	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	

HTA licensed organisation Commercial Supplier Imported (from outside England, Wales or N.Ireland)

If Yes, list lot numbers (or other identifier) & the corresponding assigned unique sample ID	Lot N°: G221222206024, G221222206017, G221222206027	Assigned Unique ID (Procuero): S00281243, S00281244
If Yes, provide Project Ref N°:	BRA060 & BRA010	
If Yes, provide the name of the PI:	Dr Rob Thomas	


A2. Details of Receipt

Date/Time of receipt	Date	07/12/2022	Time:	16:00
ID of Receiver	Name:	Jon Harriman	Dept:	CBE
ID of Supplier/Provider	Name:	Anthony Nolan	Country:	UK

PART B: To be completed by the Receiver

B1. Inspection and Quarantine

Has a biological risk assessment for the use of this material been approved?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	Ref Number: BRA060 & BRA010
Physical integrity of the material(s) acceptable?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	If No, describe action taken
Quantity received correct?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	If No, describe action taken
Labelling correct and legible?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	If No, describe action taken
Is the relevant documentation attached to this form?	C of A or equivalent evidence of quality	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
If No, add reference or details	Agreements to enable transfer of material eg MTA, SLA	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A


to ensure traceability	Details/evidence/assurance of consent	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
	Other (describe)	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Temporary storage - Quarantine location (as applicable)	Building/Room	Received 16:00 07/12/22. Stored in HTA box in lab cold room overnight. Processed 08/12/22 (H21, JH) into 2x Mixed donor 1mL vials @4.48E6. Frozen via passive cooling in H34 -80C ULT freezer until 09/12/22 and transferred to LN2 cryo-bank.
	Storage Unit ID	4Fri-HTA & Cryo-HTA
	Within storage unit location ID	Bank 7 Rack 5 Box A 24,25
	Date/Time of quarantine	09/12/2022 12:00
Submitted by:	Signature: 	Date: 07/12/22

PART C: To be completed by the departmental Quality Manager

C1. Quality Assurance Checks

Has the sample/specimen been screened for infectious biological agents?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Has the donor been screened for infectious biological agents?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Has the sample/specimen been screened and tested negative for mycoplasma?	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Is there evidence that the supplier/provider operates under national or international standards or other recognised certification?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
For HTA licensable material, is there sufficient evidence to support the requirements for storage and use of the material under the University's HTA Research Licence?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Is there sufficient evidence to support the requirements for HTA licensing exemption?	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> N/A

C2. Approval for release from quarantine

Can the material be released from quarantine and transferred to designated storage area or released for processing?	<input type="checkbox"/> Yes <input type="checkbox"/> No	
If No, provide recommendations for deposition of the material and the results of any action relating to non-conforming material.	<input type="checkbox"/> Accept as is, but with extra controls	
	<input type="checkbox"/> Rework or reprocess to meet the specified requirements	
	<input type="checkbox"/> Test to meet specified requirements	
	<input type="checkbox"/> Return to supplier/provider	
	<input type="checkbox"/> Disposal	
If Yes, provide details of storage location (as applicable)	Building/Room	
	Storage Unit ID	
	Within storage unit location ID	As above
	Database Reference	
	Date/Time of transfer	
Approved by:	Signature: 	Date: 13/12/22