PAT 003.

## Loughborough University The Centre for Biological Engineering Doc Ref: FS008.1 Version N°: 1.0 Issue Date:

ARF No: CBE/ARF/000

: HTA-PR-FORM/007

PART A: To be completed by the R	eceiver (a separate	form must be co	ompleted for	each sample t	уре)		
A1. Details of Sample/Specimen							
Type/ID: (eg primary cell, cell line, tissue, fluid, excreta, biological agent)						⊠Human □Animal	
Format / Quantity: (eg vials, slides, etc	4x50mL tube	4x50mL tube containing ~21mL ficol unit. 4 donors total					
Tissue site/Organ source:	Cord Blood	Cord Blood				□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.						⊠Yes □No	
Is the material obtained from an HTA licenced Tissue Bank with REC approval for generic research use?						⊠Yes □No	
Is the material obtained for storage and use under a project specific NHS REC approval?						□Yes ⊠No	
Is the material licensable under the HTA? Indicate source below:						⊠Yes □No	
	mmercial Supplier	☐ Imported (fro	m outside Eng	gland, Wales o	r N.Ireland)	*	
If Yes, list lot numbers (or other identifier) & the corresponding assigned unique sample ID		Lot N°: G221222205017, G2212222050 G221222205075, G2212222050			Assigned Unique ID (Procuro): S00272571, S00272572, S00272573		
If Yes, provide Project Ref N°:	BRA060 & B	BRA060 & BRA010					
If Yes, provide the name of the PI:  Dr Rob Thomas							
A2. Details of Receipt							
Date/Time of receipt	Date	19/10/2022		Time:	16:00		
ID of Receiver	Name:	Jon Harriman		Dept:	СВЕ		
ID of Supplier/Provider	Name:	Anthony Nolan		Country:	UK		
PART B: To be completed by the I	Receiver						
<b>B1.</b> Inspection and Quarantine							
Has a biological risk assessment for the use of this material been approved?		⊠Yes □No		er: BRA060 & BRA010			
Physical integrity of the material(s) acceptable?		⊠Yes □No	If No, describe action taken				
Quantity received correct?		⊠Yes □No	If No, describe action taken				
Labelling correct and legible?		⊠Yes □No	If No, describe action taken				
Is the relevant documentation attached to this form?	C of A or equivalent quality  Agreements to ena		⊠Yes □No □N/A  ⊠Yes □No □N/A				
If No, add reference or details	SLA	ansier of Mies Livo Liv/A					