Loughborough University The Centre for Biological Engineering

Acquisition and Receipt of Biological Materials

Doc Ref: FS008.1

: HTA-PR-FORM/007

Version N°:

1.0

Issue Date:

ARF No: CBE/ARF/000 47

PART A: To be completed by the	e Receiv	ver (a separate	form must be	completed for	each sample t	type)		
A1. Details of Sample/Specimen)							
Type/ID: (eg primary cell, cell line, tisst fluid, excreta, biological agent)	ue, body	CO34+ primery progenitus from perpherol blood cryo preserved vias					☑Human □Animal	
Format / Quantity: (eg vials, slides, e	etc)	3× 0	yo preserve	d vias		1		
Tissue site/Organ source:		blood				*	□N/A	
Batch N°:	,				9 9 K			
Is the sample/specimen consider If No, go to section A2.	red to b	e Relevant Ma	t Material under the Human Tissue Act (HTA)?					
Is the material obtained from an	ned from an HTA licenced Tissue Bank with REC approval for generic research use?						□Yes ☑No	
Is the material obtained for stora	s the material obtained for storage and use under a project specific NHS REC approval?						□Yes ☑No	
Is the material licensable under t	s the material licensable under the HTA? Indicate source below:						✓Yes □No	
☐HTA licensed organisation	Comme	rcial Supplier	☐ Imported (fro	om outside Eng	land, Wales o	r N.Ireland)	**	
If Yes, list lot numbers (or other identifier) & the corresponding assigned unique sample ID		Lot N°: 🦪	Lot N°: 1405082118A			Assigned Unique ID (Procuro): \$\int 00178\; 207 908 909		
			BRAOTO					
If Yes, provide Project Ref N°: If Yes, provide the name of the PI: Paf. Rob Thom			ob Thomas					
A2. Details of Receipt								
Date/Time of receipt		Date	05/03	12020	Time:	12:30)	
ID of Receiver		Name:	SON HARRE	IMAL!	Dept:	CBE		
ID of Supplier/Provider	Ÿ	Name:	Axul Bis		Country:	U.K.		
PART B: To be completed by the	Receiv	er						
B1. Inspection and Quarantine								
Has a biological risk assessment for the use of this material been approved?		use of this	✓Yes □No	Ref Number:	CBE/BRA/UIO			
Physical integrity of the material(s) acceptable?		✓Yes □No	If No, describ	e action taken				
Quantity received correct?		✓Yes □No	If No, describ	describe action taken				
Labelling correct and legible?			✓Yes □No	If No, describe action taken				
Is the relevant documentation	C of A or equivalent quality		evidence of	. □Yes □No □N/A		ria di		
attached to this form?	Agreements to enab			ØYes □No □N/A		2		
to ensure traceability		ails/evidence/assurance of		ØYes □ No I	□N/A	4	a .	

	Other (describe))	□Yes □No □N/A	u,	
Temporary storage - Quarantine location (as applicable)	Building/Room		1127 (500178907, 9 H30 (500178908) Incubator B	09)	
	Storage Unit ID		Being 7		
	Within storage (unit location ID	Botton Shelf Nerces & Box C. 6		
	Date/Time of qu	uarantine	00		
Submitted by:	Signature:		Date: 05/03/2020		
PART C: To be completed by the d	lepartmental Qu	iality Manager	这种是类型的发展		
C1. Quality Assurance Checks					
Has the sample/specimen been sci	☐Yes ☐No ☐N/A				
Has the donor been screened for in	□Yes □No □N/A				
Has the sample/specimen been scr	oplasma?	□Yes □No □N/A			
Is there evidence that the supplier or other recognised certification?	□Yes □No □N/A				
For HTA licensable material, is then and use of the material under the	□Yes □No □N/A				
Is there sufficient evidence to supp	□Yes □No □N/A				
C2. Approval for release from qua	CONTRACTOR OF THE PARTY OF THE				
Can the material be released from	quarantine and	transferred to design	nated storage area or	□Yes □No	
released for processing?	Barrier de la composition della composition dell				
released for processing?		☐Accept as is, bu	t with extra controls		
	or deposition		ocess to meet the		
If No, provide recommendations for of the material and the results of a	nny action	Rework or repr	ocess to meet the		
If No, provide recommendations for	nny action	Rework or repr	ocess to meet the nents recified requirements		
If No, provide recommendations for of the material and the results of a	nny action	☐Rework or repr specified requirer ☐Test to meet sp	ocess to meet the nents recified requirements		
If No, provide recommendations for of the material and the results of a	nny action	☐Rework or repr specified requirer ☐Test to meet sp ☐Return to supp	ocess to meet the nents recified requirements		
If No, provide recommendations for of the material and the results of a relating to non-conforming materi	any action al.	☐Rework or repr specified requirer☐Test to meet sp ☐Return to supp☐Disposal	ocess to meet the nents recified requirements	As alone	
If No, provide recommendations for of the material and the results of a	any action al.	☐Rework or repr specified requirer☐ ☐Test to meet sp☐ ☐Return to supp☐ ☐Disposal☐ Building/Room☐	ocess to meet the nents ecified requirements lier/provider	As alone.	
If No, provide recommendations for of the material and the results of a relating to non-conforming material. If Yes, provide details of storage lo	any action al.	☐Rework or represpecified requirer ☐Test to meet sp ☐Return to supp ☐Disposal ☐Building/Room ☐Storage Unit ID	ocess to meet the nents recified requirements lier/provider	As alove	
If No, provide recommendations for of the material and the results of a relating to non-conforming material. If Yes, provide details of storage lo	any action al.	☐Rework or represpecified requirer☐Test to meet sp☐Return to supp☐Disposal☐Building/Room☐Storage Unit ID☐Within storage ur	ocess to meet the nents recified requirements lier/provider	As alone.	