Loughborough University The Centre for Biological Engineering Doc Ref: FS008.1 : HTA-PR-FORM/007 Acquisition and Receipt of Biological Materials Issue Date: 18.05.2021

ARF No: CBE/ARF/00353

PART A: To be completed by the	e Receiver (a separat	te form must be	e completed f	or each samp	le type)		
A1. Details of Sample/Specimen	1						
Type/ID: (eg primary cell, cell line, tiss body fluid, excreta, biological agent)	ue, Frozen PBM	Frozen PBMC aliquots				⊠Human □Animal	
Format / Quantity: (eg vials, slides,	etc) 80* cryovial	80* cryovials					
Tissue site/Organ source:	PBMCs	PBMCs					
Batch N°:	ax3418, 341	ax3418, 3418101420K					
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?							
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	
☐HTA licensed organisation ☒	Commercial Supplier	☐ Imported (f	rom outside I	England, Wale	s or N.Ireland)	¥	
If Yes, list lot numbers (or other identifier) & the corresponding assigned unique sample ID	Lot N°: 3418101420k	(z, *	Assigned Unique ID (Procuro): S00208847 S00208848			
If Yes, provide Project Ref N°:	J13772	J13772					
If Yes, provide the name of the F	s, provide the name of the PI:				9 6		
A2. Details of Receipt	1			Ť			
Date/Time of receipt	Date	185/05/2021		Time:	10.00 am		
ID of Receiver	Name:	Maryam Shariatzadeh		Dept:	Centre for Biological Engineering		
ID of Supplier/Provider	Name:	Cambridge Bioscience, research Donors LTD		Country:	UK		
PART B: To be completed by the	e Receiver						
B1. Inspection and Quarantine							
Has a biological risk assessment for the use of this material been approved?		⊠Yes □No	Ref Number:154				
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 evidence of quality		⊠Yes □No □N/A				
	Agreements to enable transfer of material eg MTA, SLA		□Yes □No ⊠N/A				

If No, add reference or details to ensure traceability	Details/evidence	ce/assurance of		96 °
	Other (describe) □Yes □No ⊠N/A		* .	
Temporary storage - Quarantine location (as applicable)	Building/Room	/Room		
	Storage Unit ID			
	Within storage unit location ID			
	Date/Time of quarantine			
Submitted by:	Signature: Maryam Sharia	Date: 18.05.2021		
PART C: To be completed by the	e departmental O	uality Manager		
C1. Quality Assurance Checks				
Has the sample/specimen been	□Yes □No □N/A			
Has the donor been screened fo	☐Yes ☐No ☐N/A			
Has the sample/specimen been	□Yes □No □N/A			
Is there evidence that the suppli standards or other recognised or	□Yes □No □N/A			
For HTA licensable material, is the storage and use of the material	☑Yes □No □N/A			
Is there sufficient evidence to su	□Yes ☑No □N/A			
C2. Approval for release from q	uarantine			
Can the material be released fro released for processing?	m quarantine and	d transferred to de	signated storage area or	□Yes □No
If No, provide recommendations for deposition of the material and the results of any action relating to non-conforming material.		☐Accept as is, b	out with extra controls	
		☐Rework or rep specified require	process to meet the ements	
		☐Test to meet s	specified requirements	
		☐Return to sup	plier/provider	
Constitution (□Disposal		n
		Building/Room		and was
		Storage Unit ID		1000
If Yes, provide details of storage location (as applicable)		Within storage u	ınit location ID	
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- China		Date/Time of tra	nsfer	
Approved by:		Signature:	cul	Date: 18/5/2-1