MRC/Wellcome-funded Human Developmental Biology Resource (HDBR)

Registration Form

Project Title:			
Principal Investigator:			
Position Held:			
Institution:			
Address:			
Telephone No:	E	Ext No:	
E-mail address:			
am applying to use the HDI This is an extension of a pre			□ No
Project Number:	Date project-speci ethics received:	ific	
Date Project Registered:	Date MTA Agreed:	:	
Date Project Started:	Date Project Finis	hed:	

This form consists of 3 sections:

- 1. Project background and finance details.
- 2. Materials requested.
- 3. Conditions of use of the HDBR, contact details and signatures.

Section 1. Project Description

1. To allow the HDBR Steering Committee to review your project, please give a short summary (maximum of 500 words) of the background/history to your proposed research.
Please include details of:
a. Background and introduction to project.

b. Will the proposed project involve any of the following:
Create neuronal "assembloids" derived from more than one tissue type – for example neuronal-sensory cell types (e.g. eye, ear) or neuronal-motor (muscular) cell types?
Produce germ cells (including primordial germ cells)?
Culture cells or their derivatives indefinitely, or retain cells for use in further studies?
Operate a 'biobank' of HDBR-derived cellular material (e.g. organoids), to be provided to secondary users?
Transfer HDBR tissue into animals?
Use HDBR derived tissue for commercial purposes?
None of the above.
c. Objectives of proposed study.
- List the main objectives of your proposed study in order of priority.
d. Technical summary.
- Outline the techniques and experiments planned.
 If culturing tissue (cells or organoids) include details on the maximum length of time the cultures will be maintained and what will happen to the cells/tissue once the project has been completed.

e.	Justification of numbers of samples being requested.
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	 Attention should be given to the number of samples required to complete the study. Sample numbers required to produce statistically meaningful results must be considered alongside the understanding that the samples are scarce.
f.	What data will be generated and where will it be stored?
	 It is strongly encouraged by our funders to deposit all data generated in open access databases.
	 Whenever generating data on microscope slides, please record the experimental details using the Slide Experimental Details form (http://hdbr.org/uploads/default/factsheets/1529682017 f53 hdbr-slide-record.pdf). The data will be uploaded to the HDBR Atlas website (www.hdbratlas.org) - a publicly available resource for sharing human gene expression data.
	 When returning your results slides to the HDBR (see Section 3 HDBR Terms and conditions) please include a completed Slide Submission form (http://hdbr.org/uploads/default/factsheets/1529682017_f53_hdbr-slide-record.pdf).

2.	Project description suitable for lay person review (150-word summary).
	Please note this information will be included in the HDBR's annual ethics report, to be reviewed by the Health Research Authority.
3.	Project funding (ongoing or planned). Please specify:
	a. Funding body:
	b. Grant start date:
	c. Grant end date:
4.	Collaborators:
•	
	Please provide details of any project collaborators and provide full details of their part in the proposed project (continue on a separate page if there is more than one collaborator).
	a. Name:
	b. Institution:

Describe who will benefit from the proposed research.

g.

d. Telephone number:e. Email address:f. Role in project:

provide details:

g. Will HDBR tissue, its derivatives or data be shared with this collaborator? Please

5. Who	ere did you hear about us?	
a. b. c. d. e. f. g.	HDBR Website (www.hdbr.org): MRC/Wellcome Trust information: Personal recommendation: Conference: HDBR flyer: Paper citation: Other, please specify:	
con		ta generated from this study, be used for rt of a filed patent application? If yes, please

Finance Details

Upon registration of an HDBR project an initial administration fee of £500 (plus VAT) for academic users and £1,000 (plus VAT) for commercial users will be payable for each registered project. A purchase order for this payment will be requested upon registration of the project.

An annual administration fee of £500 plus VAT (academic) and £1,000 plus VAT (commercial) will be payable per annum for the duration of the project and is non-transferable upon renewal or completion of the project.

Other costs may be incurred during the project lifecycle (i.e., courier shipment fees, dry ice costs, solution/reagents, IHGES, tissue and slide processing charges, other agreed costs).

All financial transactions external to Newcastle University require the provision of a purchase order made payable to Newcastle University and a copy provided to hdbr@newcastle.ac.uk. Invoices cannot be raised, and payments cannot be allocated to projects if this document is not provided by the user.

UK Credit card payments for the £500 (plus VAT) administration fee can be made via our webstore without a purchase order. A webstore receipt will be supplied upon completion of the transaction. However, if an invoice is required, a purchase order will need to be supplied. https://webstore.ncl.ac.uk/product-catalogue/faculty-of-medical-sciences/medical-sciences/hdbr

HDBR material can only be provided once payment is received.

Please provide the following information:

Full contact details of your Finance Department:							
Postal address:							
Tel. number:							
Contact person (+ email address):							
PO Number / Grant Code (NCL internal only)							
International users:							
Bank Name and Address:							
Bank Account No:							
Sort Code:							
IBAN							
Swift Code							
VAT registration number:							•
(UK/EU Countries only)							
Commercial organisations can address	queries	to	the	Resource	Manager	at	either

hdbr@ncl.ac.uk or hdbr@ucl.ac.uk.

I confirm that I have read and understood the financial requirements outlined in the project registration form.

Name:	Signature:

Section 2. Materials Requested

In most cases, a maximum of 25 tissue samples will be released from the HDBR in the first instance. A project update form will need to be completed and returned before further tissues can be provided, and again after every 25 samples to a maximum of 100 tissues. When requesting slides, up to 20 slides may be requested from each wax block from a maximum of 25 wax blocks.

The total number of samples requested across tables 1a (archived tissue) and 1b (fresh tissue) must not exceed 100. In the case of slide requests, the wax block from which the slides were sectioned is considered to be one sample.

Table 1a: for receiving archived material (fixed, wax-embedded or frozen tissue, sectioned material on microscope slides, cDNA, RNA OR DNA) or tissues from another HDBR project (please provide Embryo ID numbers if possible).

Please complete table **1b** on the following page if you are applying to access material from the ongoing collection (fresh tissue including fresh frozen).

Source of m requested	aterials	Material	Material preparation					
Tissue or Organ	*Stage (specify CS or pcw)	Number of wax blocks	Number of slides* (maximum of 20 slides per wax block)	Number of frozen tissues	Number of genomic DNA aliquots*	No of RNA aliquots*	No of cDNA aliquots*	
		Sub total	Sub total	Sub total	Sub total	Sub total	Sub total	

Total number of tissues requested (wax blocks, frozen tissues, or cDNA/RNA/DNA* aliquots) = Total number wax blocks / slides* requested =

^{*} Charges apply for labour and consumables

^{**} Please refer to staging guides on HDBR website (http://hdbr.org/factsheets).

Table 1b: for receiving material from the ongoing collection (fresh tissue)

Please complete table **1a** on the previous page if you are applying to access archived material (fixed, wax-embedded or frozen tissue, sectioned material on microscope slides, cDNA, RNA or DNA). Please indicate the TOTAL number of samples required to complete the project.

Т	issue Requeste	ed	Sam	ple Proce	essing	Comments
Number of samples	Organ/Fetal abnormality	**Stages Please indicate CS or pcw	Fixative	Media (L15 is standard, if not please specify)	Storage temperature	(e.g. if all organs must come from same fetus, or if any special instructions are required for tissue processing)

The total number of samples requested (tables 1a and 1b) should not exceed 100. Samples will be dispatched in batches of no more than 25 following which a completed feedback form must be returned in order to receive the next batch of samples.

Total number of samples requested =

Total number of archived samples requested (table 1a) =

Total number of fresh tissues request (table 1b) =

^{*} Charges apply for labour and consumables

^{**}Please refer to staging guides on HDBR website (http://hdbr.org/factsheets).

1. Receiving tissue	from the HDBR.			
Do you wish to receive	e tissue from:			
	ollect in person [sollect in person [Arrange for colle Will arrange a c		urier 🗆
Note:				
EU: EORI Number red USA: CDC Declaratio		s: EORI Number: mport of UN3373 Categ	gory B sampl	es into the USA
Contact Details				
Please confirm the co collection of material fro are working directly on	om the HDBR. Idea	al candidates are post-c		
1st Contact:				
Name and position				
Telephone No:			Ext. No:	
E-mail address:				
2 nd Contact:				
Name and position				
Telephone No:			Ext. No:	
E-mail address:				

	Material Transfer Agreement (MTA)	
	Before any material can be released from the HDBR MTA to cover the transfer of material to your Institute required from the two HDBR centres: Newcastle (New (UCL). Once your project has been officially registered documents will be sent to your institution to agree and	ution. Separate MTAs will be reastle University) and London ed with the HDBR, draft MTA
	Tissue must not be sent to collaborators without prior of a Material Transfer Agreement being in place to cover even to researchers within the same institution (pleahttps://hdbr.s3.eu-west-2.amazonaws.com/downloads/	the movement of the material, se see operating principles –
	Name of person who will prepare MTA (usually within Contracts Dept.):	
	Position:	
	Postal Address:	
	Telephone number:	
	Email:	
4.	Project ethics approval	
4.	Project ethics approval Most UK-based HDBR projects are covered by the HD ethical approval. Confirmation of favourable ethical opi http://hdbr.org/ethical-approvals .	
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4.	Most UK-based HDBR projects are covered by the HD ethical approval. Confirmation of favourable ethical opinhttp://hdbr.org/ethical-approvals. Please specify which of the following apply: a. The project will be based solely within the Uk undertaken using the HDBR research tissue based. b. The project will be based outside the UK as	nion can be downloaded here: K and it will be ank ethics. Ind local ethics I

Section 3. Conditions of use of the HDBR

On signing this form, the applicant agrees to be bound by the following conditions, which have been established by the Joint Steering Committee, which oversees and regulates the operation of the HDBR:

I understand that HDBR samples are not screened for viruses or other pathogens and should be treated as potentially pathogenic. A comprehensive risk assessment must be performed and documented acknowledging this risk before receiving any tissue from the HDBR. Whilst some limited donor information is recorded and can be supplied to researchers, all samples are collected anonymously which means there is no possibility of returning to donors for additional clinical testing or to obtain identifiable patient data.

Additional information can be found in our factsheet "Limitations of working with human material" (http://www.hdbr.org/factsheets/).

HDBR Terms and Conditions

The following Terms govern use of HDBR material. Please read these carefully you agree to be bound by these Terms and Conditions when registering your project with the HDBR and signing below.

I agree to:

- · respect the value of this human material.
- abide by the Human Tissue Authority codes of practices <u>www.hta.gov.uk</u>, ISSCR guidelines and conditions laid out in the ethics approval of the HDBR research tissue banks.
- use the material only for the approved purpose described in my HDBR registration and shall not transfer this material (or derivatives including cell lines) to any third party without the prior written consent of the HDBR.
- submit a new project application if I wish to use HDBR material for a different purpose.
- perform a laboratory risk assessment before any HDBR material is used.
- pay the annual HDBR registration fees, and if required any IHGES, tissue and slide processing charges.
- ensure MTAs are in place to cover transfer of the material if appropriate.
- submit a project report to the HDBR every 6 months following project registration or after, 25 tissue samples have been received.
- inform the HDBR once a manuscript has been accepted for publication.
- inform the HDBR if I change institution or university.
- inform the HDBR before commercialising any results or materials provided by the HDBR.
- use the following wording in any publications arising from this work (including presentations and posters), "The human embryonic and fetal material was provided by the Joint MRC/Wellcome Trust (grant# MR/R006237/1) Human Developmental Biology Resource (www.hdbr.org)." Any subsequent secondary publications arising from data generated from HDBR material should similarly acknowledge the HDBR.
- give appropriate acknowledgement to the HDBR in any subsequent secondary publications arising from data generated from HDBR material.

I also agree to:

• the HDBR withholding material until a project report has been received.

- the HDBR refusing my application or terminating the supply of HDBR material without prior notice, and without giving a reason for this decision.
- the tissue requested being released in a staged manner depending upon satisfactory and demonstrable research progress being made.
- the project being terminated if a project report is not submitted when requested.
- the project registration continuing for a maximum of five years.
- my HDBR project title and institutional affiliation being published on the publicly accessible HDBR website www.hdbr.org/projects

Following publication, or one year after the submission of a final report to the HDBR, I agree to:

- return all gene and protein expression images taken from microscope slides.
- provide details of antibodies and probes used for all slides returned http://hdbr.org/uploads/default/factsheets/1529682017_f53_hdbr-slide-record.pdf
- HDBR capturing images from returned slides and adding them to the HDBR gene expression database www.hdbratlas.org.
- submit all sequencing data and data generated from all high through-put studies to a publicly accessible database.

Please note that publication of manuscripts in Open Access journals is strongly encouraged by our funders; for example, see www.doaj.org. Our funders Open Access policy can be found on their websites - https://wellcome.ac.uk/funding/managing-grant/open-access-policy and https://mrc.ukri.org/research/policies-and-guidance-for-researchers/open-access-policy/.

The HDBR provides human embryo/fetal tissue to all users on an equal access basis, however, I acknowledge that the HDBR, the funders and all users of the resource accept no liability for any overlap of the project aims, methods, outcomes or outputs (including publications) that arise from use of HDBR material. The HDBR policy on potentially overlapping research projects is available at https://hdbr.org/policy-on-overlapping-projects.

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☐ I do not agree to have my contact details forwarded to other researchers where my project overlaps with another HDBR project.

From time to time we would like to contact you with details of services we provide, and updates to service provision due to holidays or exceptional events (such as the COVID-19 pandemic).

☐ I agree ☐ I do not agree

Subscribe to our free Substack to receive our newsletters, updates on services and latest publications directly into your inbox. https://hdbratlas.substack.com/



Information regarding how your information will be processed can be found in our privacy notices here https://www.ncl.ac.uk/data.protection/dataprotectionpolicy/privacynotice/ and here https://www.ucl.ac.uk/legal-services/privacy/general-privacy-notice

To be completed by the Principal Investigator:

The Principal Investigator is expected to be the grant holder and will take overall responsibility for the project and ensure that the above conditions are adhered to.

I agree that I have read and understand and will abide by the HDBR terms and conditions outlined above.

Name:	
Signature:	
Date:	

Statement of Support from the Applicant's Head of Department

I have discussed the proposed project with the applicant and support his/her use of human embryonic/fetal material to be supplied by the HDBR.

Note: If the applicant is also the head of department, please ask a deputy or a person in an equivalent position to complete this section.

Name:				
Department:				
Institution:				
Address:				
Telephone No:			Ext. No:	
E-mail address:				
Signature:		Date:		

Please return the completed form to:

Dr S. Lisgo

Biosciences Institute, Newcastle University, International Centre for Life, Central Parkway, Newcastle upon Tyne. NE1 3BZ.

Email: HDBR@ncl.ac.uk

OR

Dr N. Solanky

Developmental Biology & Cancer Programme, UCL, Great Ormond Street Institute of Child Health, 30 Guilford Street, London. WC1N 1EH.

Email: HDBR@ucl.ac.uk