

INTERNAL BIOBANK CODE

AX-CaaXXX

IMP-039 rev 23 (19/12/2024)

### Highlighted fields are to be filled by the Biobank

Unless stated otherwise, all fields are compulsory. Please, consider to which area you are requesting the samples:

NEUROLOGICAL TISSUE BANK (section 4.I) – Postmortem neurological tissues В

UMOUR AND TISSUE BANK (section	on 4.II) – Biopsies from tumors and other pathologies
SIOLOGICAL FLUIDS BANK (section	4.III) – Liquid biopsies from different pathologies
1. APPLICANT INFORMATION	
	ain responsible of the project's grant)
Name & Surname	
Department/Unit	
Institution	
Sample destination institution	
Postal address	
Telephone	
E-mail	
coordinate, collaborative or multic	(Co-PI) Please fill in this section if you are collaborating within a entric project despite not being the main PI. official evidence (i.e., list of Co-IPs in the approved project).
Name & Surname	
Department/Unit	
Institution	
Sample destination institution	
Postal address	
Telephone	
E-mail	
	<u> </u>
2. PROJECT INFORMATION	
n case that your Ethics Committee	be request, the approval of your project by your Ethics Committee.  belongs to Hospital Clínic, the approval of the project and the approval of the approval o
or sample request may be processe	ed simultaneously. It is essential to contact the Biobank beforehand.
	d by your Ethics Committee contemplate in its original version the use ssociated data? And the realization of the experiments you request the
	u present an amendment to your Ethics Committee and send us its
approval.	
PROJECT TITLE	
Funding Agency / Promoter	
Official project code	
PROJECT SUMMARY (approx. 500	0 words)



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PROJECT GOALS (approx. 100 words)	
<b>EXPERIMENTS TO BE CARRIED OUT WITH THE REC</b>	QUESTED SAMPLES (approx. 100 words)
PROJECT BILLING DATA	
☐ <b>FUNDACIÓ CLÍNIC</b> ; please state grant code:	
☐ <b>OTHER</b> ; please state:	
Entity	
NIF/VAT number	
Postal address	
Contact person (if different from PI)	
Other information to add to the invoice	

It is recommended to plan the global necessity of samples to avoid subsequent applications regarding to the same project.

#### 3. PROJECT EXTENSION

### EXTENSION 'ESMENA X' () ('ESMENA' code to be filled by the Biobank)

**NOTE:** If you have previously requested samples and associated data to the Biobank for this specific project, we consider it as a PROJECT EXTENSION. In this case, you are required to, in addition to the corresponding sample section (sections 4, 5, 6), provide the following information:

REASON FOR THE PROJECT EXTENSION (approx. 100 words)
BRIEFLY DESCRIBE THE EXPERIMENTS TO BE PERFORMED WITH THE PROVIDED PROJECT EXTENSION
(approx. 100 words)

#### 4. REQUESTED SAMPLES AND/OR DATA

REQUEST () (code to be filled by the Biobank)

AMMEND X () (code to be filled by the Biobank)

**TYPE OF REQUEST:** Please specify the type of sample and/or data requested to the corresponding bank from HCB-IDIBAPS Biobank: I. Neurological Tissue Bank; II. Tumour and Tissue Bank; III. Biological Fluids Bank. Section IV applies only for data requests.

Please remember that <u>exclusive use of whole tissue blocks for research is not allowed</u>. Reference Law: RD1716/2011 (BOE-A-2011-18919). Therefore, blocks should always remain under the auspices of the Anatomy Department or Biobank core facility.



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☐ Sample request (please fill the	following sect	ions I. I	I. and/or III)		
☐ Sample & Data request (please	fill the follow	ing sec	tions I. II. And <sub>/</sub>	or III)	
☐ Data request (please fill the see	ction IV)				
I. NEUROLOGICAL TISSUE BAN	V (can be re	moved	if not applic	abla)	
	ik (call be rei	illoved	п посаррпс	abiej	
SELECTION CRITERIA  Post-mortem delay required (< ho	urc)				
Other conditions (please specif		٩٠ -			
severity, stage of pathology, age,		u.			
Select the type of neurodegenerat		d num	ber of cases th	nat are o	f your interest:
☐ Alzheimer's disease		Nº cas	ses		
☐ Amyotrophic Lateral Sclerosis		Nº cas	ses		
☐ Corticobasal degeneration		Nº cas	ses		
☐ Creutzfeldt-Jakob disease		Nº cas	ses		
☐ Frontotemporal lobar degenera	ation (please	Nº cas	ses		
specify subtype)					
☐ Huntington disease		Nº cas	ses		
☐ Lewy Body disease (please spec	cify subtype)	Nº cas	ses		
☐ Multisystemic atrophy		Nº cases			
☐ Progressive supranuclear paral	ysis	Nº cas	ses		
☐ Other (please specify) Nº cases					
Select the type of samples that are of your interest:					
☐ Fragment of frozen brain tissue	e				
☐ Histological sections from frozen brain tissue			Nº sections:		Thickness:
☐ Histological sections from cryopreserved bra			Nº sections:		Thickness:
tissue (fixed with 4%PFA 24h, and					
☐ Histological sections from para	iffin-embedde	d	Nº sections:		Thickness:
brain tissue samples	formaldobyd	,			
☐ Fragment of brain tissue in 4%		2	Nº aliquots (	600ul/ali	anot).
☐ Ventricular CSF (post-mortem)			iv- anquots (	ουσμί/ απ	quotj.
☐ Other (please specify):					
Select the areas that are of your interest:					
☐ Orbitofrontal cortex	☐ Cerebellar hemisphere			lus globe	
☐ Prefrontal cortex	☐ Dentate nucleus		☐ Thal	amus	
☐ Premotor cortex	☐ Midbrain			thalamus	
☐ Supplementary motor area	☐ Substantia nigra		☐ Luys	nucleus	
☐ Motor cortex	☐ Locus coerelus		☐ Meyı	nert nucleus	
☐ Precuneus cortex	□ Pons		☐ Hipp	ocampus	
☐ Anterior cingulate cortex ☐ Medulla oblo		blonga	ta	☐ Amy	gdala
☐ Posterior cingulate cortex ☐ Cervical sp		oinal co	rd	☐ Olfac	tory bulb
☐ Temporal cortex ☐ Thoracic spin		pinal co	ord	☐ Optio	chiasm
☐ Parietal cortex	☐ Lumbar spinal cord		rd	☐ Pituitary gland	
☐ Occipital cortex	☐ Striatum		☐ Pineal gland		
☐ Insula	☐ Caudate nucleus ☐ Other				
☐ Cerebellar vermis	☐ Putamen	nucleus	5		



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Requested data	:
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Clinical characteristics an	d/or other specification	s to consider for sample sel	ection:	
			☐ Not applicable	
		ender, age, clinical diagnosi:	s, main neuropathological	
findings and PMD), please	e mention il others are i	ieeded:	□ Not applicable	
			☐ Not applicable	
☐ This sample request requires scientific collaboration.  II. TUMOUR AND TISSUE BANK (can be removed if not applicable)				
II. TUMOUR AND HSS	OE BANK (can be rem	noved if not applicable)		
SELECTION CRITERIA				
Tissue / Organ				
Pathology				
Nº cases				
Do you require normal tis				
☐ Yes (compulsory)	☐ Yes (optional)	│	☐ Not applicable	
Clinical characteristics an	d/or other specification	s to consider for sample sel		
			☐ Not applicable	
Samples are associated w are needed:	ith basic clinical data (ge	ender, age, organ, diagnosis	), please mention if others	
			☐ Not applicable	
you need a specific nur	nber of sections.	Minimum size:		
☐ Fresh tissue		Minimum size/Nº aliquots	··	
☐ Processed tissue			·	
☐ Frozen tissue (snap-fro		Minimum size:	I	
☐ Frozen tissue (in OCT)	sections in slides	Nº slides/case:	Thickness:	
☐ Frozen tissue (in OCT)	sections in tubes	Nº tubes/case:	Thickness:	
☐ Paraffin-embedded tis	sue sections in slides	Nº slides/case:	Thickness:	
☐ Paraffin-embedded tis	sue sections in tubes	Nº tubes/case:	Thickness:	
☐ Sections of TMA (Tissu	ie Microarrays)		ne Biobank's personnel ds and sample availability.  Thickness:	
☐ Digitalised histological	preparations	Nº slides/case: Please specify staining:		
☐ Other (please specify):		1 2422 ch 22 1 220 21.		
□ This sample request rec	uires scientific collabora			
III. BIOLOGICAL FLUIDS	BANK (can be remov	ved if not applicable)		
SELECTION CRITERIA				
Pathology				
Other conditions (please required: age, gender, etc	-			



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Select the type of samples that are of your interest:

TYPE OF SAMPLES	NUMBER OF REQUESTED	SAMPLE AMOUNT / CONCENTRATION NEEDED FOR	VOLUME OF SAMPLE ALIQUOTES (plasma,
□ DNA	CASES	EXPERIMENTS (only for DNA)	serum, etc.)
☐ Plasma			
Serum			
☐ Other (please	specify):		
		linical data (gender, age, organ, diagnosis)	, please mention if others
			☐ Not applicable
	quest requires science  DATA (can be r	emoved if not applicable)	
SELECT THE BAN	NK FROM WHICH YO	OU REQUESTED THE DATA:	
☐ Neurological	tissue bank		
☐ Tumor and T	issue bank		
☐ Biological flu	ids bank		
List of the neces	ssary data for the p	roject. Please specify the number of case	es to study.

### 5. MATERIAL TRANSFER AGREEMENT (Only INTERNAL RESEARCHERS)

The use of the samples transferred hereinafter (the "MATERIAL") by the PI ("RECIPIENT") is regulated by the Spanish Law 14/2007 of Biomedical Research.

The recipient is committed to comply with the following obligations:

- To use the supplied MATERIAL exclusively for carrying out the presented project, which was previously evaluated by its relevant Ethics Committee. In the event of a substantial change in the development of the project that affects the use of the MATERIAL, the RECIPIENT must inform the BIOBANK, which will expressly decide on the authorization of the new use of the MATERIAL.
- To safeguard and ensure the traceability of the samples.
- Not to give the MATERIAL to other researchers and/or institutions who are not included in the initial PROJECT.
- To always guarantee the confidentiality of the samples and data. The commitment of confidentiality
  and limitation of use persists throughout the period in which the data are maintained, and this
  cannot be extended beyond that necessary to fulfill the research purposes indicated in the project
  and the obligations linked to it.



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- The RECIPIENT, when dealing with coded data, undertakes not to attempt to re-identify the subject.
- To assume responsibility for the proper and safe handling of the MATERIAL under appropriate biosafety conditions and by trained personnel in the RECIPIENT's laboratory to ensure appropriate risk containment.
- To inform the BIOBANK and ensure access to the corresponding data, if during the research a finding relevant for the health of the donor or his/her relatives is obtained.
- To mention the origin of the MATERIAL in all communications and scientific publications resulting from the research using the samples and/or data, with the following formulations in conjunction:
  - <u>In Materials and Methods</u>: "Samples and data from patients included in this study were provided by the HCB-IDIBAPS Biobank (B.0000575), integrated in the Platform ISCIII Biobanks and Biomodels and they were processed following standard operating procedures with the appropriate approval of the Ethics and Scientific Committees".
  - <u>In Acknowledgements</u>: "We are indebted to the HCB-IDIBAPS Biobank for sample and data procurement."
- To send a report of all published communications and scientific articles to the BIOBANK once the
  results derived from the use of the samples and/or data have been published, and to make any raw
  data of interest derived from the analyses of the MATERIAL available to future researchers who
  request the same samples.
- Upon completion of the project or termination of the contract, the RECIPIENT must DESTROY surplus samples used for said purpose as directed by said institution or RETURN them to the BIOBANK.
- To cover the expenses incurred by the BIOBANK according to a previously accepted budget, as well as shipping costs, if any, within 30 days after issuance of the invoice.
- To contract a shipping company that ensures proper transport of the MATERIAL and complies with quality standards. The BIOBANK does not assume responsibility for any damage that may occur during transport.

$\square$ By selecting this box, the applicant agrees to comply with all regulations for use of the samples
provided. This section is only for applicants within FUNDACIÓ DE RECERCA CLÍNIC BARCELONA-INSTITUT
D'INVESTIGACIONS BIOMÈDIQUES AUGUST PI I SUNYER.

#### 6. DATA MANAGEMENT ASSOCIATED TO THE REQUEST

#### 6.1 DATA PROCUREMENT TO INTERNAL RESEARCH GROUPS - Promoters of the Biobank

This section applies to Biobank sample and data procurement to **internal research groups at the Hospital Clínic de Barcelona** that are **promoters of the Biobank**. It is of upmost importance that the research group verifies that all the associated data and variables that are requested are already specified in the Data Management Plan of the approved study protocol.

Plan of the approved study protocol.
□ Commitment not to use the biological material and the data for any purpose other than the one initially indicated, except in the case that the personal data is to be reused for the purpose of health and biomedical research. In this case, an additional IC is requested for a specific purpose and to use the data for purposes related to the area in which the initial study is scientifically integrated (Art. 97 additional provision seventeen, 3/2018 of December 5, Protection of Personal Data and guarantee of digital rights), and also maintain the data confidentiality.
□ Data minimization commitment (the use of data will be appropriate, relevant, and limited)
6.2. SEARCH FOR ADDITIONAL DATA – Internal (no promoters of the Biobank) and External Research Groups
Will a search for additional data be necessary? YES □ NO □



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Data search by the Biobank			
The Biobank data management stafj require a considerable extended amo		n a routine basis. Nevertheless, some pr ncorporated ad hoc upon demand.	ojects
☐ Biobank Data Manager (name):	☐ Biobank Documentalist (	(name): Not applicable	
<u>Data search by the research group</u>			
count on a professional not related t set without providing tracking of the	o the research project aims, we code to the research team it he name of the profession	the research group, the research team swho will register the data and code each self.  Self.  Seal from the clinical group who is in cha	h data
Name:	Affiliation/Research team:	☐ Not applicable	
7. DATA FOR THE SHIPMENT (		are mentioned in the approved pro	ject.
Name & Surname			
Department/Unit			
Institution			
Postal address			
Telephone E-mail			
Courier Account Number (if aplicable)			
8. REQUEST SIGNATURE			
PRINCIPAL INVESTIGATOR (F	., 00	cable, COLLABORATOR IGATOR (CO-PI)	
Signed (Name and Surname): Date:	Signed ( Date:	(Name and Surname):	



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☐ Availability of these samples has been reviewed and approved by the HCB-IDIBAPS Biobank.

Hereby, the HCB-IDIBAPS **Biobank** approves the technical feasibility of the project and confirms the availability of:

□ All requested samples/data
 □ Some of the requested samples/data
 If so, specify which ones:
 □ None. The Biobank commits to coordinate the prospective tissue collection and create a new Biobank cohort

This approval is subject to internal committee review and approval, as well as external committee review and approval, and the Scientific Director Signature.