

IMP-039 rev 12 (15/02/2023)

1. APPLICANT INFORMATION

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

NEUROLOGICAL TISSUE BANK (section 4) – Postmortem neurological tissues TUMOUR AND TISSUE BANK (section 5) – Biopsies from tumors and other pathologies BIOLOGICAL FLUIDS BANK (section 6) – Liquid biopsies from different pathologies

	ain responsible of the project's grant)
Name & Surname	
Department/Unit	
Institution	
Postal address	
Telephone	
E-mail	
coordinate, collaborative or multio	(Co-PI) Please fill in this section if you are collaborating within a centric project despite not being the main PI. official evidence (i.e., list of Co-IPs in the approved project).
Name & Surname	onicial entactines (i.e., i.e. of each of an area approved project).
Department/Unit	
Institution	
Postal address	
Telephone	
E-mail	
2. PROJECT INFORMATION	
Does the research project approve of human biological samples? And ☐ Yes ☐ No	ed simultaneously. It is essential to contact before the Biobank. ed by your Ethics Committee contemplate in its original version the use the realization of the experiments you request the samples for? u present an amendment to your Ethics Committee and send us its
PROJECT TITLE	
- 1: 4 / 5	
Funding Agency / Promoter	
Funding Agency / Promoter Official project code	
	0 words)
Official project code	0 words)
Official project code PROJECT SUMMARY (approx. 50	
Official project code	



PROJECT BILLING DATA	
☐ FUNDACIÓ CLÍNIC; please state grant code:	
☐ OTHER; please state:	
Entity	
NIF/VAT number	
Postal address	
Contact person (if different from PI)	
Other information to add to the invoice	
3. PROJECT EXTENSION	
NOTE: If you have previously requested camples	to the Richard for this specific project, we consider it as
	to the Biobank for this specific project, we consider it as ired to, in addition to the corresponding sample section
(sections 4, 5, 6), provide the following informat	
(sections 4, 5, 6), provide the following informati	
REASON FOR THE PROJECT EXTENSION (approx	x. 100 words)
	REFORMED WITH THE PROVIDED PROJECT EXTENSION
(approx. 100 words)	
4. NEUROLOGICAL TISSUE BANK (can be re	emoved if not applicable)
SELECTION CRITERIA	
Post-mortem delay required (< hours)	
Other conditions (please specify, if require	ad.
severity, stage of pathology, age, gender, etc.)	.u.
3010.11, 310.80 0. patriology, 480, 801.40.1, 310.1,	
Select the type of neurodegenerative disease an	nd number of cases that are of your interest:
☐ Alzheimer's disease	Nº cases
☐ Amyotrophic Lateral Sclerosis	Nº cases
☐ Corticobasal degeneration	Nº cases
☐ Creutzfeldt-Jakob disease	Nº cases
☐ Frontotemporal lobar degeneration (please	Nº cases
specify subtype)	
☐ Huntington disease	Nº cases
☐ Lewy Body disease (please specify subtype)	Nº cases
☐ Multisystemic atrophy	Nº cases
☐ Progressive supranuclear paralysis	Nº cases
☐ Other (please specify)	Nº cases
Select the type of samples that are of your inter	rest:

Nº sections:

☐ Histological sections from frozen brain tissue

Thickness:



☐ Histological sections from cryopreserved brain tissue (fixed with 4%PFA 24h, and 30% sacarose 48h		Nº sections:		Thickness:
☐ Histological sections from paraffin-embedded brain tissue samples		Nº sections:		Thickness:
☐ Fragment of brain tissue in 4%	s formaldehyde			
☐ Ventricular CSF (post-mortem		Nº aliquots (500μl/ali	quot):
☐ Other (please specify):	,			
Select the areas that are of your interest:				
☐ Orbitofrontal cortex	☐ Cerebellar hem	nisphere	☐ Pallio	dus globe
☐ Prefrontal cortex	☐ Dentate nucleu	ıs	☐ Thalamus	
☐ Premotor cortex	☐ Midbrain		☐ Hypothalamus	
☐ Supplementary motor area	☐ Substantia nigr			nucleus
☐ Motor cortex	☐ Locus coerelus	-		nert nucleus
☐ Precuneus cortex	☐ Pons		☐ Hippocampus	
☐ Anterior cingulate cortex	☐ Medulla oblong	gata	☐ Amy	
☐ Posterior cingulate cortex	☐ Cervical spinal			ctory bulb
☐ Temporal cortex	☐ Thoracic spinal			chiasm
☐ Parietal cortex	☐ Lumbar spinal (tary gland
□ Occipital cortex	☐ Striatum		☐ Pinea	· -
☐ Insula	☐ Caudate nuclei	us	☐ Othe	
☐ Cerebellar vermis	☐ Putamen nucle			•
5. TUMOUR AND TISSUE BAN	K (can be remove	d if not applica	ıble)	
SELECTION CRITERIA				
Tissue / Organ				
Pathology				
Nº cases		-		
Do you require normal tissue adj			ase?	
	(optional)	□ No		☐ Not applicable
Clinical characteristics and/or oth	er specifications to	consider for san	iple sele	
				☐ Not applicable
Samples are associated with basic are needed:	c clinical data (gende	er, age, organ, di	agnosis),	please mention if others
				☐ Not applicable
Select the type of samples that are of your interest: - For slides, we normally cut at 3-5uM as standard, please state if you need another thickness. - For tubes, we have standardized a protocol depending on tissue and sample size, please let us know if you need a specific number of sections.				
☐ Fresh tissue	М	inimum size:		
☐ Frozen tissue sections in slides		№ slides/case:		
☐ Frozen tissue sections in tubes	Nº	Nº tubes/case:		
☐ Paraffin-embedded tissue sect	ions in slides Nº	Nº slides/case:		
☐ Paraffin-embedded tissue sect		№ tubes/case:		
☐ Sections of TMA (Tissue Microarrays)		º tubes/case:		
☐ Sections of TMA (Tissue Micro	arrays) De	esign agreed v		e Biobank's personnel s and sample availability.



6. BIOLOGICAL FLUIDS BANK (can be removed if not applicable)

SELECTION CRITERIA	
Pathology	
Other conditions (please specify, if	
required: age, gender, etc.)	

Select the type of samples that are of your interest:

TYPE OF SAMPLES	NUMBER OF REQUESTED CASES	SAMPLE AMOUNT / CONCENTRATION NEEDED FOR EXPERIMENTS (only for DNA)	VOLUME OF SAMPLE ALIQUOTES (plasma, serum, etc.)	
□ DNA				
☐ Plasma				
☐ Serum				
☐ Other (please, specify):				

7. 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.
- 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.



	proper transport of the MATERIAL and complies with sume responsibility for any damage that may occur					
☐ By selecting this box, the applicant agrees to comply with all regulations for use of the sample provided. This section is only for applicants within FUNDACIÓ DE RECERCA CLÍNIC BARCELONA-INSTITUTO D'INVESTIGACIONS BIOMÈDIQUES AUGUST PI I SUNYER.						
8. DATA FOR THE SHIPMENT OF SAMPLES						
	ated below are mentioned in the approved project.					
Name & Surname						
Department/Unit						
Institution						
Postal address						
Telephone						
E-mail						
Courier Account Number						
(if aplicable)						
9. REQUEST SIGNATURE						
PRINCIPAL INVESTIGATOR (PI) of the PROJECT	If applicable, COLLABORATOR INVESTIGATOR (CO-PI)					
Signed (Name and Surname): Date:	Signed (Name and Surname): Date:					