



HRCI/HRB Joint Funding Scheme 2024

PART B1: Application form

IMPORTANT INSTRUCTIONS

Please fill in the application form with reference to the Applicant Guidance Notes. These contain more detailed explanations of the type of information expected under each question.

**The final file size of the application form must be a maximum of 2MB. Supporting figures, Gantt chart, and other associated documents (e.g. signature page) must not be embedded in the main 'application' document but provided as separate files.

**Any figures to support the project description must be provided in a single additional document up to a maximum file size of 2MB.

**The Gantt chart should be provided as a separate file with a maximum file size of 2MB.

Please use font Calibri, size 11

PROJECT TITLE (maximum 20 words):

Section 1: DETAILS OF Lead AND CO-APPLICANTS

1.1 Lead Applicant details:

Name:	
Title/position:	
Department or equivalent:	
Name of Institution:	
Address:	
Tel number:	
Mobile number:	
Email address:	

1.2 Co-Applicant details:

Details of all Co-Applicants associated with this research proposal should be listed.

Note: For additional co-applicants please copy and paste table as necessary (**up to a maximum of 6 Co-Applicants can be listed**).

Co-Applicant 1	
Name:	
Title/position:	
Department or equivalent:	
Name of Institution:	
Address:	
Tel number:	
Email address:	

1.3 Host Institution for the award

For the purposes of contracting, payment, and management of the award, HRB funds can only be awarded to HRB approved Host Institutions, including those in Northern Ireland. Please note this call is open to non-island of Ireland Host Institutions. The Host Institution for the award is normally that of the Lead Applicant, but it may be another organisation/institution designated by the research team, where it is clearly justified.

Please list the Host Institution for this award, i.e. the institution to which the research award will be made. Provide details of the Dean of Research/CEO/equivalent authorised person of that institution.

A list of island of Ireland Host Institution, recognised by the HRB at the time of this call going live can be found on this [list](#).

Note: For island of Ireland Host Institutions, to be eligible to apply for funding, an Institution must be approved as a HRB Host Institution no later than two calendar months before the closing date of a call, only pre-approved Host Institutions will appear on the list linked above. Host Institutions outside of the island of Ireland do not need to be approved but must be prepared to sign up to HRB Terms & Conditions.

Name of research institution:	
Address:	
Contact person (Dean of Research/ CEO/ Equivalent authorised personnel of institution):	
Title/position:	
Tel number:	
Email address:	

Section 2: PROJECT DESCRIPTION

2.1 Project Lay Summary

Please provide a plain English summary such that it is clear, easy to understand, and is easily accessible to a **broad lay audience** (maximum **300 words**).

2.2 Project Abstract of research proposal (maximum **300 words**).

2.3 Relevance of research to strategic aims of the charity or charities (maximum **300 words**)

Please set out the relevance of your application in addressing the strategic aims of the charity or charities (in the case that two charities are co-funding) and why the charity/charities should select your application to bring forward to the HRCI/HRB-jointly nominated selection committee. Where available, refer specifically to the strategic plan of the charity/charities you apply to, and to any other relevant strategy documents (maximum **300 words**).

2.4 Keywords

Please list up to **five keywords** that specifically describe your area of research.

2.5 Project Description

The Project Description must include:

- Research Question
- Current Knowledge, Background to the Area, Relevance and Knowledge Gap
- Overall Aim
- Objectives and Deliverables (plus Gantt chart or alternative)
- Research Design and Methodological Approach
- Impact Statement
- IP Considerations
- Dissemination and Knowledge Translation Plan
- Project Management
- FAIR Data Management and Stewardship
- Public and Patient Involvement (PPI) in the Research Project
- Gender and/or Sex Issues in the Research Project

- Potential Safety Risks and Ethical Concerns
- Biobanking (where appropriate)
- Project Description Figures (where appropriate)
- References

Please ensure that your application is focused, and that sufficient evidence is provided to enable the international peer reviewers and grant selection panel to reach a considered judgement as to the quality of your research application, its significance and its feasibility.

***Any figures to support the project description must be provided in a single additional document up to a maximum file size of 2MB.**

2.5.1 Research Question (maximum 50 words).

2.5.2 Current knowledge, Background to the Area, Relevance and Knowledge Gap (maximum 1200 words).

2.5.3 Overall Aim (maximum 100 words).

Objectives and deliverables (maximum 60 words for each objective and 150 for deliverables).

Please add at least 3 individual objectives. Objectives should be SMART (specific, measurable, achievable, realistic and time-bound). For each objective please list a subset of deliverables which will be used to measure progress. Note that the stated objectives and deliverables will be used to monitor progress throughout the lifetime of the award. Timelines should be set against objectives/deliverables in your Gantt chart.

Objective 1 and associated deliverables

Objective 2 and associated deliverables

Objective 3 and associated deliverables

You must provide a **Gantt chart** which lists the above objectives and deliverables against the estimated timelines for completion, together with any additional milestones/key dates and roles and responsibilities of the team etc. Please note that the preparation and submission of Data Management Plans should also be added as deliverables/milestones of the Programme. The Gantt chart should be provided as a separate file with a **maximum file size of 2MB**.

2.5.4 Research Design and Methodological Approach (maximum 4500 words)

Please review the Applicant Guidance notes carefully for the details required in this section

Has an iteration of the proposed research been submitted to any HRB award scheme in the last 3 years?

Yes

No

(If yes) Include the award scheme and year of previous submission in your answer below. Please briefly describe the changes that have been made to the application. Have the recommendations from the previous peer, panel, or public review you received influenced the changes you have made? The word limit is **300 words**.

2.5.5 Project Management (maximum 600 words).

2.5.6 Public and Patient and Carer Involvement in the Research Project (maximum 600 words).

Important: The PPI section needs to be written as a plain English summary such that it is clear, easy to understand, and is easily accessible to a lay audience.

Are you including PPI in your application?

If Yes

Please describe all PPI at each stage of the research cycle:

- Identifying and prioritising the research question

- Design

- Conduct

- Analysis
- Oversight
- Dissemination

For each stage, please include the purpose of this involvement and where applicable how PPI has influenced/changed what work has been planned.

This section should be a succinct summary of public involvement activities. Provide information on the individuals/groups and the ways in which they will be involved. PPI contributors should be representative of the relevant people and communities impacted by the research topic. **Where members of the public or patients are involved, they should be compensated for their time and contributions; this should be reflected in the project budget.**

Please ensure to provide more detail in other sections as appropriate.

If No: please explain why PPI is not relevant to your project.

2.5.7 Gender and/or sex issues in the research project (maximum 400 words).

Are there potential sex (biological) considerations for this research?

Are there potential gender (socio-cultural) considerations for this research?

If so, outline how sex and/or gender analysis will be integrated in the design, implementation, evaluation, interpretation, and dissemination of the results of the research application.

If not, you must clearly demonstrate why it is not relevant to the research application; have you done a literature search to confirm this?

2.5.8 Impact Statement (maximum 400 words).

2.5.9 Biobanking

Does your application include an element of biobanking?

Yes

No

If Yes, you must submit a completed *Infrastructure Agreement form: Form C2* with details of the *biobank*.

If Yes, you must submit a completed Infrastructure Agreement form with details of the biobank. Please describe how you will ensure good practice for biobanking components in this project, with particular regard to quality of sample collection, processing, annotation and storage. The word limit is **400 words**.

2.5.10 Potential Safety Risks and Ethical Concerns (maximum 400 words).

2.5.11 Dissemination and Knowledge Translation Plan (maximum 500 words).

2.5.12 Outline of FAIR data management and stewardship (maximum 500 words).

2.5.13 IP Considerations (maximum 500 words).

2.5.14 References

Provide a **list of publications/references (maximum 30)** cited in the project description above

Example

Smyth, B.P. & O'Brien, M. (2004) Children Attending Addiction Treatment Services in county Dublin, 1990-1999. *European Addiction Research*, 10(7455) pp. 68-74.

2.6 Project Description Figures

Any figures to support your Project Description can be included as a single additional upload. A **maximum of 5 figures**, which can be a combination of images, graphs, tables, scales, instruments, or surveys, can be included as a single document. They must not be embedded within the text of the Project Description. Additional references should not be included here. The maximum size is **2MB**. Files should be doc, docx, or pdf.

Section 3: DETAILS OF RESEARCH TEAM

Research Team Roles

3.1 Lead Applicant Role

Outline the role of the PI in the project on a day-to-day basis including amount of time to be spent working on the project either as a percentage or proportion of a full time equivalent (FTE) (maximum **250 words**).

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3.2 Co-Applicants Role

For each Co-Applicant, please identify the type of Co-Applicant they are here (Researcher Co-applicant, Knowledge User Co-applicant, or PPI Co-applicant) and outline their role in the project on a day-to-day basis including amount of time to be spent working on the project either as a percentage or proportion of a full time equivalent (FTE). Describe the specific contribution and responsibilities of the Co-Applicant (maximum **250 words**).

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3.3 Collaborator's Role

For each Collaborator, please outline their role in the project on a day-to-day basis including amount of time to be spent working on the project either as a percentage or proportion of a full time equivalent (FTE).

Note: For each collaborator a signed **Collaboration Agreement Form** must be provided. A template Collaboration Agreement Form is available with all application forms from the HRCI-registered research charity (maximum **250 words per Collaborator**).

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3.4 Personnel

Give full details of all personnel to be funded through this project. Please fill in the following table for each person. If more tables are required please copy and paste as necessary.

Specify Personnel Type	
State percent time on project	
State specific role in the project	
Person known – y/n	
If yes, name	
If yes, address	
If yes, Present position	
If yes, Academic and Professional Qualifications	
Give a detailed justification for the nature of the research personnel relative to the scale and complexity of the project. The word limit is 400 words .	

SECTION 4: RESEARCH INSTITUTION INFRASTRUCTURE AND SUPPORT

4.1 Host Institution Infrastructure and Support

Describe the infrastructure, facilities, specialist expertise and other support available at the Host Institution and/or at other sites where the research will be conducted. Please include details of critical supports in areas such as statistics, methods, trial management or regulatory expertise where this is being provided above and beyond the activities/expertise of members of the research team. (maximum **400 words**).

4.2 Access to Research Infrastructure

Do you plan to avail of the advice, research design, data management services and/or other forms of support from a Clinical Research Facility/Centre (CRF/CRC), other infrastructure unit (e.g., Centre for Applied Medical Imaging, CAMI, Centre for Support and Training in Analysis and Research (CSTAR)) at research design or implementation stages?

Yes

No

If **Yes**, the following information must be provided (**maximum 400 words**):

- Name and address of the facility/centre/network
- Information on the nature and stage/s of the input/advice/collaboration/service;
- Rationale for the choice of facility/centre/network
- How the proposed involvement enables the planned research to be undertaken to the required quality or timescale.

If **No**, applications involving patients which do not detail such input, advice and/or support (and where this expertise is not clearly evident within the applicant team) should justify why they have chosen not to access such support (**maximum 400 words**).

Where applicable a signed **Infrastructure Agreement Form** must be provided. Failure to provide an Infrastructure Agreement Form(s) will result in the application being deemed ineligible. Electronic signatures are acceptable.

Section 5: PROJECT DURATION AND BUDGET

5.1 Project duration and budget total

Please indicate the expected length of the proposed project in months, proposed start date and the total budget requested. The minimum duration is 12 months and the maximum is 36 months.

The maximum total value of an award is **€300,000**. There is no set limit per annum: costs should be allocated in the year expected to occur.

Duration:
Budget Total:
Proposed start date (not earlier than 01 October 2024):

5.2 Project Budget

See the Instructions to Applicants for detailed guidance regarding the project budget.

A **full detailed breakdown** of **costings** and **justification for all funding** is required for items listed under each subheading. You are strongly advised to seek guidance from the research office/finance office in the Host Institution before completing this section of the form. HRCI/HRB will not provide additional funding in the case of either under-estimates or over expenditure.

Use **Table 1** to provide a summary of the costs requested and **Table 2** to justify each amount requested.

Table 1: Total direct costs related to the proposal

Please provide details of the **total amount of funding** requested for each year of the research proposal (Direct Costs only). Note that HRCI/HRB awards will be up to a maximum total award value (direct costs) of **€300,000** for projects from 12 months up to 36 months.

Only include direct costs in this application. HRB will apply a rate of 30% of Total Direct Modified Costs (TDMC excludes student fees, equipment, and capital building costs) for **laboratory or clinically based research** and 25% of Total Direct Modified Costs for **desk-based research** on the HRB portion of research funding at time of contract for successful applications.

Cost Item	Year 1	Year 2	Year 3	Total
1. Personnel Costs				
a) Gross Salary (inclusive of employees' pension contribution)				
b) Employer's PRSI				
c) Employer Pension				

Contribution				
d) Student Stipend				
e) Student Fees				
2. Running Costs				
3. PPI Costs				
4. Equipment				
5. Dissemination Costs				
6. FAIR Data Management Costs				
Total Costs				

Table 2: Justification of costs.

Under each of the headings please **itemise each cost** and provide a brief but explicit **justification of the costs** claimed.

For Personnel Costs, please state the pay scale used and the level and point on the scale. This should be justified accordingly. For appointment of Research Fellows or Senior Research Fellows evidence of position must be provided at point of award.

Line Item	Justification
1. Personnel Costs a) Gross Salary b) PRSI c) Pension (max <u>200 words</u>)	
1d) Student Stipend (max <u>100 words</u>)	
1e) Student Fees (max <u>100 words</u>)	
2. Running Costs (max <u>400 words</u>)	
3. PPI Costs (max <u>200 words</u>)	
4. Equipment (max <u>200 words</u>)	
5. Dissemination Costs (max <u>200 words</u>)	

6. FAIR Data Management Costs (max 200 words)	
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5.3 Other Funding (maximum 200 words).

Please indicate if you have submitted this, or a similar application, to another HRB scheme or funding body. If this application has been submitted elsewhere, please indicate which HRB scheme or funding body, project title, result of submission or when outcome is expected and the amount of award. The word limit is 200 words.

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Failure to disclose accurately or fully will result in your application being deemed ineligible and withdrawn without further review.

Section 6: ETHICAL APPROVAL AND APPROVALS FOR USE OF ANIMALS

Ethical approval is required for all research work funded by the HRB that involves human participants, human material (including tissue) or animals. Applicants are responsible for ensuring that all necessary approvals are in place prior to the start of the research.

Applicants should allow sufficient time to obtain ethical and/or competent authority approval and/or animal licenses as a copy of such approvals must be submitted to the HRB before the initiation of the award. It is suggested that these are sought in parallel to the submission of the application to the HRB.

Please complete the following Approvals Declaration.

HRB GRANT – APPROVALS DECLARATION

Name of Principal Investigator	«Lead Applicant»
Grant Reference	«Grant Reference»
Title of Grant	«Grant Title»
Duration of Grant	«Grant Duration months» months
Commencement Date of Research	«Grant Start Date»

(A) Do you require any of the following approvals* for some/all aspects of your research programme?	Insert Yes/No in all cases
Research Ethics Committee (REC) Approval	
HPRA Authorisations for use of animals in research	
Clinical Trial Approval from HPRA	
HR-CDC Consent Declaration	

**Refers to requirements that were declared at the outset of the project and any additional requirements or amendments since then, including renewals required within the timeframe of the project/programme.*

IF Yes to any of the above, Complete (B) to (D)

(B)	Approval required from: (e.g. month 1)	Approval anticipated by: (date)	From (e.g. Committee name)
REC Approval			
Animal Authorisations			HPRA
Clinical Trial Approval			HPRA
HR-CDC Consent Declaration			HR-CDC

(C) I confirm that I have received the following	Confirm by completing details below
Research Ethics Committee (REC) Approval	<i>Date received; reference number</i>
Animal Authorisations	<i>Dates received; reference numbers</i>
Clinical Trial Approval from HPRA	<i>Date received; reference number</i>
HR-CDC Consent Declaration from HR-CDC	<i>Date received; reference number</i>

(D) I am not in a position to include confirmation of *(tick those that apply)*

REC Approval ___ Animal authorisations ___ Clinical Trial Approval ___ HR-CDC declaration ___

at this time but I hereby confirm that I will not proceed with any element of the research programme which requires approval before receipt of approval and sending an updated approvals declaration to the HRB confirming same.

PI Signature		Date	
HI Signature		Date	

Section 7 : LEAD APPLICANT AND CO-APPLICANT CVs AND COLLABORATOR PROFILES

The templates below **must** be used for CVs.

Lead Applicant and Co-Applicant CVs can be a maximum of 5 pages and should be broken down as follows: *Section 1 (max 2 pages) + Section 2 (max 1 page) + Section 3 (max 1 pages)*

LEAD APPLICANT

7.1 Lead Applicant CV

Section 1 – Required Details (max 2 pages)

Name and contact details

Please underline the name of the PI on each publication listed.							
Total Publications #							
Senior author publications #							
Journal Articles #							
Reviews #							
Book Chapters #							
Books #							
Conference associated publications* #							
Other #							
<i>*Conference associated publications can be classified into peer reviewed conference papers and edited conference proceedings where appropriate as per discipline</i>							
Details of research funding most relevant to this application as Principal Investigator/Co-Applicant (up to 5).							
<i>Please note that this section should only include funding obtained <u>as Principal or Co-Investigator</u> (expand columns as required).</i>							
<i>Start mm/yyyy</i>	<i>Duratio n (months)</i>	<i>Total amount (currency)</i>	<i>Name of funder</i>	<i>Funding Body ref. no.</i>	<i>Type (project/fellowship /other)</i>	<i>Title</i>	<i>Role of applicant</i>

7.1.2 – Supervisory experience (max 1 page)

Supervisory experience
<i>If you are planning to supervise a higher degree/postgraduate student, as part of this application, please include a brief summary of your supervisory experience to date. Please state the number of students supervised, those successfully completed and indicate how many of these are still in progress. The word limit is 200 words.</i>

7.1.3 Additional evidence for Lead Applicant

Additional evidence of experience and expertise relevant to this application

Please describe any additional experience or expertise that will provide evidence of the ability of the Lead Applicant to successfully lead the proposed project. Please use this opportunity to describe any career gaps in your CV. The word limit is **400 words**.

CO-APPLICANTS

7.3 Co-Applicant CVs

Please fill in the following table for each co-applicant. If more tables are required please copy and paste as necessary. (max. **3 pages per Co-Applicant**). Up to 6 Co-Applicants may be included on this application.

Co-Applicant 1

Section 1 – Required Details (max 2 pages)

Name and contact details			
<i>Title</i>		<i>Institution</i>	
<i>Forename(s)</i>		<i>Address Line 1</i>	
<i>Surname</i>		<i>Address Line 2</i>	
<i>Email Address</i>		<i>City/Town</i>	
<i>Position</i>		<i>County</i>	
<i>Department</i>		<i>Phone No.</i>	
Perspective as a Co-Applicant			
<i>If a Co-Applicant contributes from more than one perspective please select the dominant role</i>			
<input type="checkbox"/> Researcher Co-Applicant <input type="checkbox"/> Knowledge User Co-Applicant <input type="checkbox"/> PPI Contributor Co-Applicant			

In addition please complete one of the following sections depending on the type of Co-Applicant:

7.2.1 Researcher Co-Applicant

IF RESEARCHER CO-APPLICANT; complete the following		
ORCID (if known)		
For more information and to register please see https://orcid.org/		
ORCID iD number:		
Publications (5 most relevant)		
<i>Please fill in the table including the total number of publications and categorise that number according to the additional categories below. Please list the 5 publications that are most relevant to this application.</i>		
Date of Publication	Title	Authors

Please underline the name of the PI on each publication listed.							
Total Publications #							
Senior author publications #							
Journal Articles #							
Reviews #							
Book Chapters #							
Books #							
Conference associated publications* #							
Other #							
<i>*Conference associated publications can be classified into peer reviewed conference papers and edited conference proceedings where appropriate as per discipline</i>							
Details of research funding most relevant to this application as Principal Investigator/Co-Applicant (up to 5).							
<i>Please note that this section should only include funding obtained <u>as</u> Principal or Co-Investigator.</i>							
<i>Start mm/yyyy</i>	<i>Duration (months)</i>	<i>Total amount (currency)</i>	<i>Name of funder</i>	<i>Funding Body ref. no.</i>	<i>Type (project/fel lowship/ot her)</i>	<i>Title</i>	<i>Role of applicant</i>
Permanent Position YES <input type="checkbox"/> /NO <input type="checkbox"/> Contract Position YES <input type="checkbox"/> /NO <input type="checkbox"/> If yes, state contract end date: _____							
<p>Research Institution Letter of Support must be provided for (1) all Lead Applicants in a contract position and (2) Co-Applicants in a contract position who are seeking their own salary. The formal letter on headed notepaper and signed by the Head of School/Research Centre/Hospital must include the following information:</p> <p>[Research Institution – insert name] which is the research institution of [applicant - insert name] confirms that [applicant - insert name]: (i) holds an employment contract which extends until [insert date] or will be recognised by the research institution upon receipt of the HRCI/HRB award as a contract researcher; (ii) has an independent office and research space/facilities for which he/she is fully responsible for at least the duration of the award, and (iii) has the capability and authority to mentor and supervise the research team.</p>							
Supervisory experience							
<p>If you are planning to supervise a higher degree/postgraduate student, as part of this application, please include a brief summary of your supervisory experience to date. Please state the number of students supervised, those successfully completed and indicate how many of these are still in progress. The word limit is 400 words.</p>							

7.2.2 Knowledge User Co-Applicant

IF KNOWLEDGE USER CO-APPLICANT; complete the following
Evidence of expertise and experience in influencing decision making within knowledge user organisation(s)
<p>Knowledge User Co-Applicants should highlight their previous and current roles in influencing decision-making processes within their organization or other relevant organisations. They should also use this space to highlight their specific experiences and expertise for the Knowledge User Co-Applicant role in relation to the proposed research. The word limit is 300 words.</p> <p>A knowledge user is defined as one in a position of authority to influence and/or make data decisions about health policy or the delivery of services and that can act to ensure that the findings of the research will be translated to influence decision making and change within their (or other) organisations.</p>
Additional evidence of experience and expertise relevant to this application
<p>You may wish to include here any additional experience or expertise that will support the application. For example, you may wish to include any relevant research experience/expertise, previous experience of working in collaboration or links with researchers to produce research or evidence for health, evidence of Patient Public Involvement in your knowledge user role, and roles/responsibilities as a constructive and effective change agent. The word limit is 300 words.</p>

7.2.3 PPI Contributor Co-Applicants

IF PPI CONTRIBUTOR CO-APPLICANT complete the following
PPI Co-Applicant experience and expertise relevant to this application
PPI Co-Applicants should provide some information regarding their experience and expertise

relevant to this application. For example, they may wish to include relevant experience as a service user or carer, relevant experience from their personal lives, prior experience in PPI or any other useful background information. The word limit is **400 words**.

COLLABORATORS

7.3 Collaborators Profile

Please fill in the following table for each collaborator associated with the project. If more tables are required please copy and paste as necessary (max. **2 pages per Collaborator**). Up to 10 collaborators can be added to this application.

Not all sections will be relevant to all Collaborators.

Collaborator 1

Name and contact details							
<i>Title</i>		<i>Institution</i>					
<i>Forename(s)</i>		<i>Address Line 1</i>					
<i>Surname</i>		<i>Address Line 2</i>					
<i>Email Address</i>		<i>City/Town</i>					
<i>Position</i>		<i>County</i>					
<i>Department</i>		<i>Phone No.</i>					
Education							
<i>From</i>	<i>To</i>	<i>Qualification</i>	<i>Subject</i>	<i>Country</i>	<i>Institution</i>	<i>Class</i>	<i>Dept.</i>
Previous positions							
<i>From</i>	<i>To</i>	<i>Position</i>	<i>Organisation</i>				
ORCID							
ORCID iD							

number:							
Membership of professional body/council							
<i>Details:</i>							
Publications (5 most relevant)							
<i>Please fill in the table including the total number of publications and categorise that number according to the additional categories below. Please list the 5 publications that are most relevant to this application.</i>							
Date of Publication	Title	Authors					
<i>Please underline the name of the PI on each publication listed.</i>							
Total Publications #							
Senior author publications #							
Journal Articles #							
Reviews #							
Book Chapters #							
Books #							
Conference associated publications* #							
Other #							
<i>*Conference associated publications can be classified into peer reviewed conference papers and edited conference proceedings where appropriate as per discipline</i>							
Details of research funding most relevant to this application as Principal/Co- investigator (up to 5)							
<i>Please note that this section should only include funding obtained <u>as Principal or Co-Investigator.</u></i>							
Start mm/yyyy	Duration (months)	Total amount (currency)	Name of funder	Funding Body ref. no.	Type (project/fellowship/other)	Title	Role of applicant

Note: For each collaborator a signed **Collaboration Agreement Form** must be provided. A template Collaboration Agreement Form is available in **Part C1**. Failure to provide Collaboration Agreement Form(s) will result in the application being deemed ineligible.

All applications for funding must provide a signature page which has been signed by the Lead Applicant and the Dean of Research/CEO/equivalent authorised personnel of the Research Institution. **All signatures must be originals. Electronic versions of signatures are not acceptable.**

Checklist for submission

For all applications

Document Number	Title	Included?
B1	Application form	
Upload attachment	Gantt chart	
Upload attachment	Figures	
D1	Lead Applicant Signature page	
D2	Host Institution Signature page	

Where applicable

Document Number	Title	Included?
C1	Collaboration Agreement Form	
C2	Infrastructure Agreement Form	
C3	Letters of support	