

**INSTRUCTIONS FOR SCHOOL RESEARCH ETHICS COMMITTEE (SREC) ON THE USE OF THIS TEMPLATE**

1. As agreed by the Open Research Integrity and Ethics Committee, this Ethics Review Application Proforma is a Cardiff University template that may be modified as set out below.
2. Where [square brackets] appear in the template, the SREC must insert the required text or delete if not relevant.
3. Where comment boxes appear, these contain guidance for the SREC to consider. The comment boxes must be deleted before the Application Proforma is issued for use. Please also refer to the SREC FAQs document for guidance in specific FAQ areas.
4. The SREC is permitted to add additional information to the template (to reflect local processes) **provided** the additions do not conflict with the existing template content or other Cardiff University policies and procedures. Deletions and amendments to content within this template is generally not permitted, unless otherwise indicated. The template content represents the University's minimum expectations for the information to be provided by Researchers within an application.

**School of History, Archaeology and Religion Research Ethics Committee**

**APPLICATION FOR ETHICAL REVIEW**

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| --- | --- |
| For Office Use Only | |
| SREC Reference: [x] | Meeting/Review Date: [x] |

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|  | | | **SECTION 1. GENERAL INFORMATION** | | | | | | | | |
| Application Type: | | | | |  | £ Staff £ PGR student  £ PGT/Masters Student £ Undergraduate | | | | | |
| Research Project Title: | | | | |  |  | | | | | |
| Short Title (where applicable): | | | | |  |  | | | | | |
|  | | | For Staff Projects | | | | | | | | |
| Name of Chief/Principal Investigator: | | | | |  |  | | | | | |
| Contact details: | | | | |  |  | | | | | |
| Other members of research team: | | | | |  |  | | | | | |
|  | | | For Student Projects | | | | | | | | |
| Name of Student: | | | | |  |  | | | | | |
| Contact details: | | | | |  |  | | | | | |
| Name of Supervisor(s): | | | | |  |  | | | | | |
| Contact details: | | | | |  |  | | | | | |
| Other members of research team: | | | | |  |  | | | | | |
|  | | | **SECTION 2. SCREENING QUESTIONS** | | | | | | | | |
|  | |  | | | | | | | **Yes** |  | **No** |
| 2.1 | | Is the research project categorised as ‘Research’ (as defined in the Cardiff University [Policy on the Ethical Conduct of Research involving Human Participants, Human Material or Human Data](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/research-ethics) (Ethics Policy))?  *If* ***no*** *(i.e. the research project is a Service Evaluation or Audit), the Committee is not required to conduct a review of the proposal but may choose to do so.* ***Please contact Dr Gerwin Strobl to seek advice before proceeding with this application.*** | | | | | | |  |  |  |
| 2.2 | | Does the research project involve human participants, human material or human data (as defined in the [Ethics Policy](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/research-ethics))?  *If* ***no****, you are not required to submit the research proposal to this Committee.* ***Please do not continue with this application.*** | | | | | | |  |  |  |
| 2.3 | | Does the research project require review by an external ethics committee (refer to Appendix 1 of the [Ethics Policy](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/research-ethics))? Please note that this includes all research projects involving participants who lack the capacity to consent.  *If* ***yes****, the research project should be submitted to the relevant external ethics committee for review and does not fall within the remit of this Committee. Please contact the* [*Research Governance Team*](mailto:resgov@cardiff.ac.uk) *for further advice.* ***Please do not continue with this application.*** | | | | | | |  |  |  |
| 2.4 | | Has the research project been ethically reviewed by another university or research institution (for example, where the Chief/Principal Investigator for the research project is based at another institution)?  *If* ***yes,*** *please provide evidence of the review conducted (such as an outcome letter or communication) and the ethical review policy of the relevant institution or committee.* ***Please do not continue with this application.*** | | | | | | |  |  |  |
| 2.5 | | Is the research project exempt from ethical review under the University’s framework for the ‘[Ethical Review of Research using Secondary Data and/or Publicly Available information only’](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/research-ethics)? This framework allows certain research projects using only secondary data and/or publicly available information to proceed without ethical review by a SREC provided certain conditions are met.  *If* ***yes****,* you are not required to submit the research proposal to this Committee. **Please do not continue with this application**. If in doubt, please seek advice from Dr Gerwin Strobl. | | | | | | |  |  |  |
| 2.6 | | Does the research project fall within the scope of the [UK Policy Framework for Health and Social Care Research](https://www.hra.nhs.uk/planning-and-improving-research/policies-standards-legislation/uk-policy-framework-health-social-care-research/)? This Framework broadly applies to research taking place within, or involving, the health and social care systems.  *If* ***yes****, you will need to apply to the* [*Research Governance Team*](mailto:resgov@cardiff.ac.uk) *for Sponsorship using the Advanced Project Information Proforma (APIP) (available on the Cardiff University intranet)*. *The Research Governance Team will advise you on the approvals that are required for the research project after it has conducted a review of the APIP and supporting documentation.* ***Please do not continue with this application until you have sought advice from the Research Governance Team.*** | | | | | | |  |  |  |
| 2.7 | | Does the research project involve the collection or use of Human Tissue (including, but not limited to, blood, saliva and bodily waste fluids)?  *If* ***yes****, the research project should be submitted to the* [*Human Tissue Act Compliance Team*](mailto:hta@cardiff.ac.uk) *(HTACT) prior to submission to an ethics committee.* ***Please note that submission of a research protocol/proposal to HTACT is essential for all projects involving the collection or use of Human Tissue. Please do not continue with this application until you have sought advice from HTACT.*** | | | | | | |  |  |  |
| 2.8 | | Does the research project fall within the scope of the University’s [Security-sensitive Research Policy](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/security-sensitive-research)? This Policy broadly applies to research involving terrorism, extremism or radicalisation (or access to materials of such a nature).  *If* ***yes****, you must register the research in accordance with the Policy and comply with the IT and security arrangements contained in the Policy.* | | | | | | |  |  |  |
| 2.9 | | Has the research project received appropriate peer/scientific review? (For student research projects, review by the research project supervisor is an acceptable form of scientific review)  *If* ***no****, please obtain appropriate peer/scientific review before submitting the application to this Committee.* | | | | | | |  |  |  |
| 2.10 | | Have you and all other Cardiff University co-applicants/ /Supervisors/Members of the research team (as listed in Section 1) completed the University’s [Research Integrity Online Training Programme](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/training/research-integrity-online-training-programme) and any other required training?  *If* ***no****, you must complete the training before submitting the application to this Committee.* | | | | | | |  |  |  |
| 2.11 | | Has your supervisor both read and approved this form? | | | | | | |  |  |  |
|  | | | If the research project involves the use of animals, please contact the Cardiff University Biological Standards Office [bso@cardiff.ac.uk](mailto:bso@cardiff.ac.uk) to seek further advice. | | | | | | | | |
|  | | | **SECTION 3. PROJECT SUMMARY** | | | | | | | | |
| 3.1 | |  | | Summarise the research project (including the purpose and its methodology) using language that would be understood by a lay person. | | | | | | | |
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| 3.2 | |  | | State the research question(s). | | | | | | | |
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| 3.3 | |  | | Estimated start date. | | | | | | | |
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| 3.4 | |  | | Estimated end date (usually the end of data collection). | | | | | | | |
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| 3.5 | |  | | Is the research project funded? *If* ***yes****, please name the funding body.* | | | | | | | |
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| 3.6 | |  | | Are there any potential conflicts of interest? *If* ***yes****, please confirm the action you propose to take to address such conflicts.*  *Information and guidance on conflicts of interest is contained in the* [*Research Integrity Online Training Programme*](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/training/research-integrity-online-training-programme) *and the* [*Research Integrity and Governance Code of Practice*](https://www.cardiff.ac.uk/research/our-research-environment/integrity-and-ethics/research-integrity-and-governance)*.* | | | | | | | |
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| 3.7 | |  | | Does the research project **only** involve the use of common methodologies previously approved by the School of History, Archaeology and Religion Research Ethics Committee? *If* ***yes****, please provide details in ‘Section 10: Supporting Documents’ below and attach the relevant documentation (e.g. protocol or standard operating procedure for the common methodologies) to this application.* | | | | | | | |
|  | | | **SECTION 4. FULL REVIEW CRITERIA** | | | | | | | | |
|  | | | Your answers to the questions in this Section 4 will help the Committee determine whether your project requires full or proportionate review.  If all ‘No’ boxes apply, your project may be considered for proportionate review.  If a 'Yes' box applies, your project will proceed to full review unless the School has approved a Standard Operating Procedure (common methodologies) for that particular criterion.  Please tick ‘Yes – CM’ if you plan to use common methodologies (CM). | | | | | | | | |
|  | | | | | | | | | **Yes** | **Yes- CM** | **No** |
| 4.1 | | Will the research project be performed without the participants’ prior consent? | | | | | | |  |  |  |
| 4.2 | | Does the research design include an element of deception, including covert research? | | | | | | |  |  |  |
| 4.3 | | Will the research project involve children under the age of 18 or ‘at risk’ (vulnerable) adults or groups?  *The* [*Cardiff University Safeguarding Children and Adults at Risk: Policy*](https://www.cardiff.ac.uk/public-information/policies-and-procedures/safeguarding) *and* [*Guidance*](https://www.cardiff.ac.uk/public-information/policies-and-procedures/safeguarding/activity-specific-guidance) *sets out examples of ‘at risk’ or ‘vulnerable’ adults.* | | | | | | |  |  |  |
| 4.4 | | Does the research project include topics which may be considered highly sensitive for participants?  *This includes sexual behaviour, illegal activities, political, religious or spiritual beliefs, race or ethnicity, experience of violence, abuse or exploitation, and mental health.* | | | | | | |  |  |  |
| 4.5 | | Does the research project require access to records of a sensitive or confidential nature, including Special Category Data?  *Special Category Data is defined in data protection legislation and currently includes information about an individual's: racial or ethnic origin; political opinions; religious beliefs; trade union membership; physical or mental health; sexual life or orientation; commission of offences or alleged offences; genetic data; and biometric data.* | | | | | | |  |  |  |
| 4.6 | | Is permission of a gatekeeper required for initial or continued access to participants?  *This includes participants in custody and care settings, or research in communities where access to research participants is not possible without the permission of another adult, such as another family member or a community leader.* | | | | | | |  |  |  |
| 4.7 | | Does the research project involve intrusive or invasive procedures?  *This includes the administration of substances, vigorous physical exercise, procedures involving pain or more than mild discomfort to participants (including the risk of psychological distress, discomfort or anxiety to participants).* | | | | | | |  |  |  |
| 4.8 | | Does the research project involve visual or audio recordings where participants may be identified? | | | | | | |  |  |  |
| 4.9 | | Does the research project involve the collection or use of human tissue (collected from the living or where less than 100 years have passed since the persons death)? | | | | | | |  |  |  |
| 4.10 | | Does the research project involve any ancient human remains?  If yes:  Have you secured the appropriate permission, if required, to excavate, export and/or sample any ancient human remains?  If applicable, have you discussed and agreed preferred options for the disposal of human remains after excavation and analysis?  Have you read and understood the School’s policy on Human Remains? | | | | | | |  |  |  |
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| 4.11 | | Does the research project involve more than a minimal risk of harm to the safety and wellbeing of participants and/or the Researchers?  *Please answer this question based on your assessment of the risks involved in this project. Further information about possible harm or potential risks to participants/researchers must be provided in Section 7 of this form.* | | | | | | |  |  |  |
| 4.12 | | Does the research project involve fieldwork?  If so:   * Have you secured the appropriate permission from the tenant and landowner? * Does the research take place outside of the UK?   If “Yes” have you gained appropriate permissions?   * Does the area of research include any Scheduled Monuments?   If “Yes” have you gained permission from the appropriate authority?   * Is the area of research special environmental interest or value (e.g., is it an SSSI)?   If “Yes” have you gained permission from the appropriate authority?   * Have you contacted the local Site and Monuments Officer? * Is there an agreement in place with the legal owner of any materials recovered regarding the deposition of material culture and archive?   If the material is to be deposited with another institution in due course, is this agreement in place? | | | | | | |  |  |  |
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| 4.13 | | Does the research involve the production of genetic data, whether human, floral or faunal?  If genetic data is to be produced:   * Is the material from outside the UK? * Has a data management and dissemination plan been developed for genetic data?   *If the DNA has not already been extracted from human tissue collected from the living, or where less than 100 years have passed since the persons death, please submit the application for review by the Human Tissue Act Compliance Team before submission to the SREC.* | | | | | | |  |  |  |
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| For each response given as ‘Yes – CM’ please provide here the name of the common methodology/methodologies being used (e.g. ‘Best Practice Guidelines for Human Remains Store’). Please also attach a copy of the common methodology/methodologies you propose to use in your project. | | | | | | | | | | | |
|  | | | **SECTION 5. PARTICIPATION AND RECRUITMENT** | | | | | | | | |
| 5.1 | | How will you identify and recruit participants to the research project? | | | | | | | | | |
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| 5.2 | | How many participants are you aiming to recruit? *If applicable, please include a breakdown of participants by type and number.* | | | | | | | | | |
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| 5.3 | | What are the inclusion and exclusion criteria for participants? | | | | | | | | | |
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| 5.4 | | Will the research project involve participants that are Cardiff University staff or students or clients of the University (or the place in which you may otherwise work)? *If applicable, please provide details.* | | | | | | | | | |
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| 5.5 | | How will the research project address recruitment of participants who are not fluent in the English/Welsh language? | | | | | | | | | |
|  | | | **SECTION 6. CONSENT PROCEDURES** | | | | | | | | |
| 6.1 | | Will informed consent be obtained from participants? If so, how? *Please include who will be taking consent, how consent will be recorded, when participants will be provided with information about the research project, and how long potential participants will be given to decide whether to take part.* | | | | | | | | | |
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| 6.2 | | Will participants be offered any incentives to take part in the research project? | | | | | | | | | |
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| 6.3 | | If a questionnaire is to be used, will you give participants the option of omitting questions they do not wish to answer? | | | | | | | | | |
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| 6.4 | | Will participants be informed that their participation is voluntary and that they may withdraw at any time and for any reason? | | | | | | | | | |
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|  | | | **SECTION 7. POSSIBLE HARM TO PARTICIPANTS/RESEARCHERS** | | | | | | | | |
| 7.1 | | Is there is a risk of the participants experiencing physical, emotional or psychological harm or distress*? If yes, please provide details of how ethical issues will be handled and how any risks will be minimised.* *Please consider whether the research project includes topics which could be considered as highly sensitive for participants.* | | | | | | | | | |
|  | | |  | | | | | | | | |
| 7.2 | | Is there a risk of the Researcher(s) experiencingphysical, emotional or psychological harm or distress? *If yes, please provide details of how ethical issues will be handled and how any risks will be minimised.* | | | | | | | | | |
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|  | | | **SECTION 8. DATA MANAGEMENT, CONFIDENTIALITY AND DATA PROTECTION** | | | | | | | | |
| 8.1 | | How, and by whom, will data be collected? | | | | | | | | | |
|  | | |  | | | | | | | | |
| 8.2 | | Will you be accessing or collecting Personal Data (identifiable personal information) as part of the research project? *If yes, please confirm what data will be accessed and/or collected (including details of the information participants are asked to provide on a written consent form) and by who.*  *Note: If your project involves Personal Data, you are advised to review the University’s* [*GDPR Guidance for Researchers*](https://intranet.cardiff.ac.uk/intranet/students/documents/Guide-to-GDPR-and-Research.pdf) *and to check whether your project requires, or would benefit from, the completion of a* [*Data Protection Impact Assessment*](https://intranet.cardiff.ac.uk/staff/supporting-your-work/manage-use-and-protect-data/data-protection/data-protection-impact-assessments) *(DPIA). It is not the role of the SREC to review or advise on DPIA’s, but if you have completed one, please confirm this below. For further advice, please refer to the ‘DPIA’ intranet page or contact* [*complianceandrisk@cardiff.ac.uk*](mailto:complianceandrisk@cardiff.ac.uk)  *A researcher* ***must*** *complete a DPIA if they answer 'yes' to* ***at least two*** *of the screening questions found* [*here*](https://intranet.cardiff.ac.uk/intranet/staff/documents/20190923-Data_Protection_Impact_Assessment_Template-CU-.docx)*. Completion of a DPIA is recommended where the researcher answers 'yes' to one screening question.* | | | | | | | | | |
|  | | |  | | | | | | | | |
| 8.3 | | How long will you retain the Personal Data collected in connection with the research project? Please also explain any data deletion arrangements.  *Note: Research records and data must be retained for the period specified in Section 2.9 ('Research Project Conduct') of the University's Research Records Retention Schedule. If identifiable information is being collected, researchers must ensure that this is limited to the information necessary to achieve the relevant purpose (data minimisation). The University expects raw data containing identifiable information (questionnaires and audio tapes for example) to be retained for the full retention period unless: (1) the identifiable information is not required to support the research or to demonstrate good research conduct;* ***and*** *(2) stringent measures have been taken to verify and ensure the integrity of any anonymised or pseudonymised records/data produced from the raw data. Where (1)* ***and*** *(2) apply, the researcher* ***must*** *take the necessary steps to remove the personal data. Consent Forms must be retained for the full retention period.*  *Please note that where UG and PGT projects do not contribute to a publication or wider research project, research records and data may be held for a shorter period. Please refer to the guidance notes in Section 2.9 of the University’s Research Records Retention Schedule for further detail.* | | | | | | | | | |
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| 8.4 | | What efforts will be made to anonymise the data collected (where possible)? | | | | | | | | | |
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| 8.5 | | Are you proposing to utilise ‘public task’ as the lawful basis for processing Personal Data for the purposes of the research project (as recommended in the University’s [GDPR Guidance for Researchers](https://intranet.cardiff.ac.uk/intranet/students/documents/Guide-to-GDPR-and-Research.pdf))?  *If no, please explain why and what alternative lawful basis you propose to use.* | | | | | | | | | |
|  | | |  | | | | | | | | |
| 8.6 | | Have you utilised/incorporated into your Participant Information Sheet the following sections from the University's template Participant Information Sheet: *'What will happen to my Personal Data'* and *'What happens to the data at the end of the research project?'*  *If* ***no****, please explain why this has not been used and how you have otherwise ensured that the relevant data protection/privacy information has been provided to participants.* | | | | | | | | | |
|  | | |  | | | | | | | | |
| 8.7 | | For how long will the collected anonymised data be retained? Please also explain any data deletion arrangements.  *Note: Anonymised research data must be retained for the period specified in Section 2.9 ('Research Project Conduct') of the University's Research Records Retention Schedule. Please note that where UG and PGT projects* ***do not*** *contribute to a publication or wider research project, research records and data may be held for a shorter period. Please refer to the guidance notes in Section 2.9 of the University's Research Records Retention Schedule for further detail.* | | | | | | | | | |
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| 8.8 | | Who will have access to the data? | | | | | | | | | |
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| 8.9 | | Will the data be shared in any way, for example through deposit in a data repository, with third parties, or a transcription service? | | | | | | | | | |
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|  | | | **SECTION 9. OTHER ETHICAL CONSIDERATIONS** | | | | | | | | |
|  | | | Please outline any other ethical considerations raised by the research project and how you intend to address these. You are obliged to bring to the attention of the SREC any ethical issues not covered in this Ethics Review Application Proforma. | | | | | | | | |
|  | | | **SECTION 10. SUPPORTING DOCUMENTS** | | | | | | | | |
|  | | | I have attached the following documents in support of this application. | | | | | | | | |
|  | | | | | | | **Yes** | **No** |  | **Version no.** (if applicable) | |
| 1 | Research Project Protocol/Proposal | | | | | |  |  |  |  | |
| 2 | Recruitment Adverts/Invitation Letters | | | | | |  |  |  |  | |
| 3 | Participant Information Sheet | | | | | |  |  |  |  | |
| 4 | Consent Form | | | | | |  |  |  |  | |
| 5 | Data Collection Tool(s) (e.g. questionnaire(s)) **or**  a detailed description of the proposed tool which provides the SREC with clear information about the parameters of the tool i.e. what themes/areas will be covered and what will be excluded. The final version of whatever Data Collection Tool is used (e.g. questionnaire) must be submitted to the SREC for approval before it is used. | | | | | |  |  |  |  | |
| 6 | Other participant communications (e.g. debrief sheets) | | | | | |  |  |  |  | |
| 7 | Evidence of Research Integrity training completion (and any other required training if relevant)? | | | | | |  |  |  |  | |
| 8 | *Applicant to list any additional documents provided to the SREC, particularly any additional documents relevant to recruitment, consent and participation or common methodologies used.* | | | | | |  |  |  |  | |
|  | | | If you have selected ‘No’ for any of the documents listed above, please confirm why these have not been provided. If a listed document is not relevant to your project, please confirm this below. | | | | | | | | |
|  | | | **SECTION 11. SIGNATURES AND DECLARATIONS** | | | | | | | | |
|  | | | **Applicant declaration**  I confirm that:   1. The information in this form is accurate to the best of my knowledge and belief and I take full responsibility for it. 2. I have the necessary skills, training and or/expertise to conduct the research project as proposed. 3. I am familiar with the University’s health and safety requirements and policies and that all relevant health and safety measures have been taken into account for the research project. 4. I am familiar with, and will comply with, the University’s [Policy on the Ethical Conduct of Research involving Human Participants, Human Material or Human Data](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/research-ethics) and the University’s [Research Integrity and Governance Code of Practice](https://www.cardiff.ac.uk/research/our-research-environment/integrity-and-ethics/research-integrity-and-governance). 5. The relevant equality and diversity considerations have been taken into account when designing the research project. 6. If the research project is approved, I undertake to adhere to the research project protocol, the terms of the full application as approved and any conditions set out by the Committee and any other body required to review and/or approve the research project. 7. I will notify the Committee and all other review bodies of substantial amendments to the protocol or the terms of the approved application, and to seek a favourable opinion from the Committee before implementing the amendment. | | | | | | | | |
|  | | | Signed: | | | | | | | | |
|  | | | Print name: | | | | | | | | |
|  | | | Date: | | | | | | | | |
|  | | | **SUPERVISOR DECLARATION (FOR STUDENT PROJECTS)** | | | | | | | | |
|  | | | I confirm that:   1. I am familiar with the University’s [Policy on the Ethical Conduct of Research involving Human Participants, Human Material or Human Data](https://intranet.cardiff.ac.uk/students/study/postgraduate-research-support/integrity-and-governance/research-ethics) and the University’s [Research Integrity and Governance Code of Practice](https://www.cardiff.ac.uk/research/our-research-environment/integrity-and-ethics/research-integrity-and-governance). 2. I have reviewed this application, and all supporting documents, and I am satisfied that the project as proposed meets the University’s ethical standards. 3. I have the necessary skills, training and or/expertise to offer appropriate supervision and support to the student researcher/applicant. 4. I will encourage the student to discuss with me, and reflect on, any ethical issues that arise during or after the project and, where relevant, I will ensure such issues are notified to the SREC. | | | | | | | | |
|  | | | Signed: | | | | | | | | |
|  | | | Print name: | | | | | | | | |
|  | | | Date: | | | | | | | | |

**Please submit the completed application and supporting documents to SHARE Committees (**[**share-committees@cardiff.ac.uk**](mailto:share-committees@cardiff.ac.uk)**), copied to Dr Gerwin Strobl (**[**StroblG@caridff.ac.uk**](mailto:StroblG@caridff.ac.uk)**).**

**Your electronic submission should contain wet-ink or electronic signatures of all relevant parties. Please note that if any information is missing, the application may be returned to you.**