**Public Benefit and Privacy Panel for Health and Social Care**

**Application Form**

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| --- | --- | --- | --- |
| **Application Control**  *Applicants should not complete the “submitted date” field* | | | |
| Application Coordinator | This is version one (v1) | | |
| Application Number |  | Submitted Date |  |
| Applicant Name |  | | |
| Proposal Name |  | | |
| Project End Date |  | | |

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| **Pre-submission checklist**  *Applicants should not fill out this section –* ***to be completed by the eDRIS coordinator*** | |
| Approved Information Governance Training | Approved training complete and certificates received  Approved training complete and certificates pending |
| Use of recognised safe haven | Yes  No |
| NHSCR Involvement | Yes  Reference number:................  Email Confirmation of approval supplied:  No |
| Is project covered by National Safe Haven generic ethics approval? | Yes  No |

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| **Supporting Documents** |
| Please list *only* supporting documents which you have clearly referenced in your application – the name of each should clearly indicate what the document/file/reference is about. |

**Note to Applicants**

Prior to completing your application form you should:

* Contact the eDRIS Team, who will assist you - [Nss.edris@nhs.net](mailto:Nss.edris@nhs.net) or by phone on 0131 275 7333
* Read and understand the separate Guidance for Applicants

Your application should be typed, not handwritten. Your eDRIS application coordinator will inform you of how to submit your application form and any supporting evidence. Before submitting your completed application, you should ensure that:

* All relevant sections of the application are complete
* Relevant supporting evidence is attached
* Individuals named on the form have read and approved its submission

Please note that submitted applications may be circulated to panel members, administrative colleagues, NHSScotland information governance and information security colleagues, Caldicott Guardians, the CHI Advisory Group and, where appropriate, non-NHS Scotland colleagues from a variety of participating partner bodies, in the course of processing. You must make your eDRIS application coordinator aware of any confidential or sensitive information contained in your application which you would consider inappropriate for circulation in such a manner. Your application could be subject to disclosure or partial disclosure under the Freedom of Information (Scotland) Act, and will be retained in line with NHSScotland information policy.

## Section 1 – People

|  |  |  |  |
| --- | --- | --- | --- |
| **1.1** | **Applicant** *Please read section 1.1 of the guidance* | | |
| **1.1.01** | Full Name: | |  |
| **1.1.02** | Title: | |  |
| **1.1.03** | Position (if PhD researcher, please also complete section 1.2): | |  |
| **1.1.04** | Professional Registration No.: | |  |
| **1.1.05** | Organisation Name: | |  |
| **1.1.06** | Address (incl. postcode): | |  |
| **1.1.07** | Email: | |  |
| **1.1.08** | Do you have an NHS contract/honorary contract? | | Choose an item. |
| **1.1.09** | Provide details of the most recent information governance training undertaken - a list of training courses is included at Appendix A of guidance notes | | |
|  | Name and institution of course: |  | |
|  | Date completed: |  | |

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| **1.2** | **PhD Supervisor** *Please read section 1.2 of the guidance* | | |
| **1.2.01** | Full Name: | |  |
| **1.2.02** | Title: | |  |
| **1.2.03** | Position: | |  |
| **1.2.04** | Professional Registration No.: | |  |
| **1.2.05** | Organisation Name: | |  |
| **1.2.06** | Address (incl. postcode): | |  |
| **1.2.07** | Email: | |  |
| **1.2.08** | Does this person have an NHS contract/honorary contract? | | Choose an item. |
| **1.2.09** | Provide details of the most recent information governance training undertaken - a list of training courses is included at Appendix A of guidance notes | | |
|  | Name and institution of course: |  | |
|  | Date completed: |  | |

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| **1.3** | **Clinical Sponsor/Lead** *Please read section 1.3 of the guidance* | | |
| **1.3.01** | Full Name: | |  |
| **1.3.02** | Title: | |  |
| **1.3.03** | Position: | |  |
| **1.3.04** | Professional Registration No.: | |  |
| **1.3.05** | Organisation Name: | |  |
| **1.3.06** | Address (incl. postcode): | |  |
| **1.3.07** | Email: | |  |
| **1.3.08** | Does this person have an NHS contract/honorary contract? | | Choose an item. |
| **1.3.09** | Provide details of the most recent information governance training undertaken - a list of training courses is included at Appendix A of guidance notes | | |
|  | Name and institution of course: |  | |
|  | Date completed: |  | |

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| --- | --- | --- | --- |
| **1.4** | **Information/Data Custodian** *Please read section 1.4 of the guidance* | | |
| **1.4.01** | Full Name: | |  |
| **1.4.02** | Title: | |  |
| **1.4.03** | Position: | |  |
| **1.4.04** | Professional Registration No.: | |  |
| **1.4.05** | Organisation Name: | |  |
| **1.4.06** | Address (incl. postcode): | |  |
| **1.4.07** | Email: | |  |
| **1.4.08** | Does this person have an NHS contract/honorary contract? | | Choose an item. |
| **1.4.09** | Provide details of the most recent information governance training undertaken - a list of training courses is included at Appendix A of guidance notes | | |
|  | Name and institution of course: |  | |
|  | Date completed: |  | |

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| **1.5 Others with access to identifiable or potentially identifiable data** *Please read section 1.5 of the guidance* | | | |
| **1.5.01** | Full Name: | |  |
| **1.5.02** | Title: | |  |
| **1.5.03** | Position: | |  |
| **1.5.04** | Professional Registration No.: | |  |
| **1.5.05** | Organisation Name: | |  |
| **1.5.06** | Address (incl. postcode): | |  |
| **1.5.07** | Email: | |  |
| **1.5.08** | Does this person have an NHS contract/honorary contract? | | Choose an item. |
| **1.5.09** | Provide details of the most recent information governance training undertaken - a list of training courses is included at Appendix A of guidance notes | | |
|  | Name and institution of course: |  | |
|  | Date completed: |  | |

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| **1.6 Others** *Please read section 1.6 of the guidance* | | | |
| *Complete this section if applicable – for each additional person* | | | |
| Full Name: |  | Involvement in Proposal: |  |
| Organisation: |  | Position: |  |

**Section 2 – Organisations & Bodies**

|  |  |  |
| --- | --- | --- |
| **2.1** | **Organisation or Body Leading Proposal** *Please read section 2.1 of the guidance* | |
| **2.1.01** | Organisation or Body Name:  *If the organisation here is an NHSScotland board, note this and go directly to question 2.1.03* |  |
| **2.1.02** | Is this a commercial organisation or body? | Choose an item. |
| **2.1.02a** | If ‘Yes’, please provide a full explanation of the organisation or body’s activity and industry sector, including any previous experience of using NHSScotland data - append supporting documentation as appropriate |  |
| **2.1.03** | Is this organisation or body wholly funding or paying for the costs of conducting the proposal? | Choose an item. |

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| **2.2** | **Main Contact for Lead Organisation** *Please read section 2.2 of the guidance* | |
| **2.2.01** | Full Name: |  |
| **2.2.02** | Title: |  |
| **2.2.03** | Position: |  |
| **2.2.04** | Email: |  |

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| **2.3** | **Organisation or Body Funding Proposal** *Please read section 2.3 of the guidance* | |
| *Complete the following section if you answered ‘No’ to question 2.1.03* | | |
| **2.3.01** | Organisation or Body Name:  *If the organisation here is an NHSScotland board note this and, go directly to section 2.4* |  |
| **2.3.02** | Is this organisation or body a commercial organisation? | Choose an item. |
| **2.3.02a** | If ‘Yes’,please provide a full explanation of the organisation or body’s activity and industry sector, including any previous experience of using NHSScotland data - append supporting documentation as appropriate |  |

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| **2.4 Other Relevant Organisations or Bodies** *Please read section 2.3 of the guidance* | | |
| *Complete this section if applicable* | | |
| Organisation Name | Nature of Business/Sector | Nature of interest in proposal |
|  |  |  |

## Section 3 – Overview

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| --- | --- | --- | --- | --- |
| **3.1** | **Proposal Essentials** *Please read section 3.1 of the guidance* | | | |
| **3.1.01** | Please specify the proposal end date | |  | |
| **3.1.02** | Is this proposal:   * an extension * a renewal of an existing approval * related to a previous application (approved or not)   Please provide details, include the reference number of the original application, and summarise the changes requested | | |  |
| **3.1.03** | Does this proposal require updates of information or to be repeated at regular intervals? If yes please advise the frequency | | |  |
| **3.1.04** | What is the substantive purpose of the proposal? (please choose **one** option from below that best matches your proposal) | | | |
|  | Patient Care | Research | | |
|  | Audit | Performance Monitoring/Management | | |
|  | Service Planning/Improvement | Health/Social Care Administration | | |
|  | Systems Implementation/Testing | Training/Education | | |
|  | Other |  | | |
|  | If other clearly defined purpose, please give details: | | | |
| **3.1.05** | Access is being requested to data from which sources? (tick as many as are relevant) | | | |
|  | A single NHS Scotland Board (excluding NSS) including any system/database  NHS National Services Scotland  More than one NHS Scotland Board including any system/database  Community Health Index (CHI) database  NHS Central Registry  Other | | | |
|  | If other, please give details: | | | |
|  | | | |
| **3.1.06** | Provide a clear and concise ***lay*** outline of the proposal (max. 250 words). This may be published on the PBPP website. | | | |
|  | | | |
| **3.1.07** | Provide a description of the aims and objectives of the proposal. | | | |
|  | | | |
| **3.1.08** | Provide a description of the envisaged benefits to the public and/or patients. | | | |
|  | | | |
| **3.1.09** | Provide a concise description of: the research study design (sample size, inclusion/exclusion criteria, time period); data collection; data processing or other means required to achieve the aims of your proposal. | | | |
|  | | | |
| **3.1.10** | Provide a clear and concise outline of any statistical methods that will be used in the project (if applicable). | | | |
|  | | | |
| **3.1.11** | Provide a diagram/description to illustrate the data flow or data linkage process envisaged (if applicable). | | | |
|  | | | |
| **3.1.12** | Does the proposal have implications for, or target, vulnerable populations? Please give details. Definitions of vulnerable populations are given in section 5 of Appendix A of the guidance notes. | | | |
|  | | | |
| **3.1.13** | Does the proposal seek access to highly sensitive data? Please give details. Definitions of sensitive data are given in section 6 of Appendix A of the guidance notes. | | | |
|  | | | |
| **3.1.14** | Does the proposal seek to use information exclusively about deceased persons? Please give details. | | | |
|  | | | |
| **3.1.15** | Describe how you have included public input / lay representation in your proposal design. | | | |
|  | | | |
| **3.1.16** | Describe any peer review undertaken, with details (for example formal review by a peer organisation or funding body, informal internal review, and review by a third party). | | | |
|  | | | |
| **3.1.17** | Describe how the proposal has been designed to demonstrate that privacy risk has been adequately assessed, is appropriately managed, and has been reduced to acceptably low levels (e.g. has a data protection impact assessment (DPIA) been carried out, if appropriate). Please provide any relevant supporting documentation. | | | |
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| **3.1.18** | Is there *any* commercial aspect or dimension to the proposal or its outcomes? If yes, please give details. | | | |
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| **3.2** | **Statutory and Regulatory Context** *Please read section 3.2 of the guidance* |
| **3.2.01** | Does your proposal have a statutory or regulatory justification - is the proposal responding to a statutory or regulatory instruction, duty or order? Please give details. |
|  |
| **3.2.02** | If your organisation will be processing personal and/or special category data as part of this proposal then please cite the lawful basis for processing under current data protection law. |
|  |
| **3.2.03** | Are there any existing information sharing agreements or contracts in place which support your proposal? Please give details and attach as supporting documentation. |
|  |
| **3.2.04** | Are regulatory approvals from outside Scotland pending or received? Please give details. |
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| **3.3** | **Research and Ethics Governance** *Please read section 3.3 of the guidance* | |
| **3.3.01** | Has your proposal sought NHS or university research ethics approval? | Choose an item. |
| **3.3.01a** | If yes, provide committee details, status of approval (i.e. pending, approved, etc) and reference number. Please attach as supporting documentation if available | |
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| **3.3.01b** | If no, explain why NHS or university research/ethics approval is not sought: | |
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| **3.4** | **Safe Havens** *Please read section 3.4 of the guidance* | |
| **3.4.01** | Do you intend to access the data requested exclusively through a safe haven listed at Appendix A of guidance notes? Please provide details of which safe haven/s.  *If you have answered ‘Yes’ you do not need to complete sections 5.1 or 5.2* | |
|  | |
| **3.4.02** | If you applying to use NHS NSS data and you do not intend to do this through the National Safe Haven, please explain why then proceed to Section 4. | |
|  | |
| **3.4.03** | Will you be accessing the safe haven remotely? | Choose an item. |
| **3.4.04** | How and at what location will you be accessing the safe haven? E.g. on a university-provided laptop from a university office. | |
|  | |

**Section 4 – Data & Data Subjects**

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| **4.1 New Data yet to be collected** *Please read section 4.1 of the guidance* | |
| Dataset/source Name | Collection by (whom)? |
|  |  |
|  |  |
|  |  |

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| --- | --- | --- |
| **4.2 All Other Existing Datasets / sources** *Please read section 4.2 of the guidance*  **Please note that contact should be established as early in the process as possible with NHS Scotland boards/Data providers to discuss data provisioning requirements for any of the applicable sources listed below.** | | |
| Dataset/source Name | Data Controller (Organisation)  **For existing dataset/sources for which the data controller is not an NHSScotland board, please append evidence of the data controllers permission to use the data** |
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| **4.2.01** | How were individuals originally informed of the use of their data? You should ensure that you include an appropriate explanation for each of the data sources which you have listed above. |
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| **4.3 Data Variables** *Please read section 4.3 of the guidance* | | | |
| Dataset/source Name | Variable | Time Period/Range | Please check to indicate if this item is used for processing only and will not be part of the output |
|  |  |  |  |
|  |  |  |  |
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| **‘**Data protection law requires that the use of either directly or indirectly identifiable data variables is minimised to those which are strictly necessary. This is known as the ‘data minimisation’ principle. In the table below please justify the need for all of the identifiable or potentially identifiable variables included in your proposal: | |
| Identifying or Potentially identifying Variable | Justification |
|  |  |
|  |  |
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| **4.4** | **Methodology** *Please read section 4.4 of the guidance* | | |
| **4.4.01** | Does the proposal require any of the following: | | |
| Data linking  Use of matched controls  Single anonymised data extract  Other (please specify): | | |
| **4.4.02** | If the proposal requires data linkage, who is undertaking the linkage e.g. eDRIS team, local analysts etc..? | | |
| **4.4.03** | What variables will be processed for linkage? | | |
| CHI Number | Forename | Surname |
| Date of Birth | Address | NHS Number |
| Postcode | Other Please Specify: | |

|  |  |  |
| --- | --- | --- |
| **4.5** | **NRS/NHSCR Data Sources** *Please read section 4.5 of the guidance* | |
| *Complete this section if access to NHSCR is required, or if there is any National Records of Scotland involvement* | | |
| **4.5.01** | Does the proposal require access to NHS Central Registry as a sampling frame for cohorts? | Choose an item. |
| **4.5.02** | Does the proposal involve flagging of individuals on the NHSCR for long term follow up? | Choose an item. |
| **4.5.03** | If yes,is flagging necessary: | |
|  | To trace and contact individuals throughout the UK? | |
|  | To be informed of fact and cause of death? | |
|  | To be informed of the incidence of on-going anonymised cancers registrations? | |
|  | To be informed of emigrations prospectively and retrospectively? | |
| **4.5.04** | Is any other NRS/NHSCR involvement required? Please provide details | |

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **4.6** | **Making Contact with Individuals** *Please read section 4.6 of the guidance* | | | | |
| **4.6.01** | Is any direct contact with any group of individuals required? If Yes, please provide details below | | | | Choose an item. |
|  | Contact Group and Method of contact | | | | Contact by (whom) |
|  | Hospital Consultants | Letter | Phone | Other (specify) : |  |
|  | Other NHSS Staff | Letter | Phone | Other (specify) : |  |
|  | General Practitioners | Letter | Phone | Other (specify) : |  |
|  | Patients/Public | Letter | Phone | Other (specify) : |  |
|  | Relatives of participants | Letter | Phone | Other (specify): |  |
|  | Others (please specify): | Letter | Phone | Other (specify) : |  |
| **4.6.02** | Please explain why contact is being made – append copies of relevant correspondence as supporting evidence | | | | |
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| **4.7** | **Community Health Index (CHI) Database** *Please read section 4.7 of the guidance* |
| *Complete this section if access to CHI Database is required* | |
| **4.7.01** | What monitoring and audit of the use of CHI is planned? Please provide details |
|  |
| **4.7.02** | What technical method will be used to access CHI (online read-only, download, other extract, anonymised extract, etc)? Please provide details |
|  |
| **4.7.03** | Have any risks been identified in the proposal which relate specifically to CHI? |
|  |

**Section 5 – Data Processing**

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| --- | --- | --- |
| **5.1** | **Access** *Please read section 5.1 of the guidance* | |
| *Complete the following section if you answered ‘No’ to question 3.4.1* | | |
| **5.1.01** | At what location is identifiable or potentially identifiable data being accessed? | |
|  | |
| **5.1.02** | Please provide details of security policies/procedures governing access to this physical and technical environment. Please append supporting documentation referencing appropriate sections. | |
|  | |
| **5.1.03** | Does this policy/procedure cover password policy in detail? Please provide details/ append supporting documentation referencing appropriate sections. | |
|  | |
| **5.1.04** | Does this policy/procedure cover user account management, including review or removal of access to sensitive/personal data, in detail? Please provide details/ append supporting documentation referencing appropriate sections. | |
|  | |
| **5.1.05** | Will individuals with access to data have individual or shared accounts? | |
|  | |
| **5.1.06** | Will the data be accessed by staff working off site e.g. staff working from home at any time during the duration of the proposal? | Choose an item. |
| **5.1.06a** | If yes, are policies/procedures in place to facilitate, monitor and audit this access? Please provide details/ append supporting documentation. | |
|  | |
| **5.1.07** | Provide any additional detail of how data is protected from unauthorised access | |
|  | |

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| **5.2** | **Storage & Use** *Please read section 5.2 of the guidance* | |
| *Complete the following section if you answered ‘No’ to question 3.4.1* | | |
| **5.2.01** | Where is data being stored and used? (location, organisation, address – refer to addresses in previous sections if appropriate) | |
|  | |
| **5.2.02** | ISO 27001 Cert. No. |  |
| **5.2.03** | Please provide details of security policy/procedure governing storage and use of data within this physical and technical environment – append supporting documentation referencing appropriate sections | |
|  | |
| **5.2.04** | Does this policy/procedure cover the implementation of up-to-date controls for the detection and prevention of malware? Please provide details/ append supporting documentation | |
|  | |
| **5.2.05** | Does this policy/procedure cover access control and auditing of system administrator activity? Please provide details/ append supporting documentation referencing appropriate sections | |
|  | |
| **5.2.06** | Does this policy/procedure cover the production of backups and the controls in place around these? Please provide details/ append supporting documentation | |
|  | |
| **5.2.07** | Does this policy/procedure describe the controls in place to prohibit unauthorised copying of data? Please provide details/ append supporting documentation referencing appropriate sections | |
|  | |
| **5.2.08** | Does this policy/procedure describe physical and site controls? Please provide details/ append supporting documentation referencing appropriate sections | |
| **5.2.09** | Does this policy/procedure cover hardware repair, replacement or disposal and protection of data from inappropriate access during such procedures? Please provide details/ append supporting documentation | |
|  | |
| **5.2.10** | Describe the systems, software and security used to store and use data - please provide details/ append supporting documentation | |
|  | |
| **5.2.11** | Is outsourced IT in use? If yes, please give details | |
|  | |
| ***Please repeat section 5.2 above for each relevant location in the proposal – see guidance*** | | |

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| **5.3** | **Transfer** *Please read section 5.3 of the guidance* | |
| **5.3.01** | Please provide details of security policy/procedure to ensure that data will be transferred in such a way that it is protected from inappropriate or unauthorised access (mention email encryption, secure file transfer protocols SFTP, device encryption, physical controls, etc, as appropriate) - append supporting documentation | |
|  | |
| **5.3.02** | At what intervals/ trigger points will data transfer take place? E.g. one off transfer, monthly intervals | |
|  | |
| **5.3.03** | Will any identifiable or potentially identifiable data be transferred outside of the UK? | Choose an item. |
| **5.3.03a** | If yes,please provide details of the country of destination, the method of transfer, the proposed location and method of storage outside of the UK, and details of any further onward transfer | |
|  | |
| **5.3.04** | Other than initial transfers from source systems, is there any copying of data required within the proposal? If yes, please give details | |
|  | |

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| **5.4** | **Dissemination** *Please read section 5.4 of the guidance* | |
| **5.4.01** | Will proposal findings be published or disseminated beyond those listed in Section 1? (*If you have answered ‘No’, go directly to section 5.5)* | Choose an item. |
| **5.4.01a** | If yes, how will proposal findings be published or disseminated, to what audience and in what format? Please give details | |
|  | |
| **5.4.01b** | If yes, what steps will be taken to ensure that persons cannot be identified in published Please give details and confirm what disclosure control policy will be applied | |
|  | |
| **5.4.01c** | If yes, are there any circumstances where a living or dead individual would be cited? (E.g. where a person consented to their data being used as a case study)? Please give details | |
|  | |
| **5.4.01d** | If yes, were any permissions to publish data required or sought (for example from data controllers)? Please provide details | |
|  | |

|  |  |
| --- | --- |
| **5.5** | **Retain/Dispose** *Please read section 5.5 of the guidance* |
| **5.5.01** | Which information/data/records retention policy will you be applying to the proposal data (details of the policy and the organisation to which it belongs)? |
|  |
| **5.5.02** | How long do you intend to retain identifiable or potentially identifiable data after the conclusion of the proposal (including archive/backup copies)? |
|  |
| **5.5.03** | Who will retain the data and where? |
|  |
| **5.5.04** | What is the purpose for retaining the data for the specified time? |
|  |
| **5.5.05** | What method of disposal or destruction will be used when this period has expired (including archive/backup copies)? |
|  |
| **5.5.06** | What evidence will be obtained that destruction has occurred (eg IT supplier certificate of destruction, etc)? |
|  |

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| --- | --- |
| **5.6** | **Review** *Please read section 5.6 of the guidance* |
| **5.6.01** | Describe how the mechanisms which safeguard data security will be audited and reviewed at regular intervals to ensure their continued efficacy |
|  |
| **5.6.02** | Describe any resource implications to any of the proposed measures for the protection of physical or technical security of information which are unresolved at the time of this application? (for example encryption of devices is an intention not yet fulfilled, training is not yet undertaken, etc) |
|  |
| **5.6.03** | Describe the breach reporting mechanisms to be invoked in the event of any inappropriate access to data or other information security incident |
|  |

## 

## Section 6 – Declaration

* I DECLARE THAT this application is accurate, and that, should it be successful, any health data made accessible will be used for no other purpose, and in no other way, than as described above.
* I UNDERTAKE TO notify the Public Benefit and Privacy Panel (PBPP) of any future changes to the purpose or manner in which data is processed in accordance with this application.
* I UNDERSTAND THAT any future applications by me, or my employing or sponsoring organisation, may be refused should any health data made accessible be used for any other purpose or in any other way than that described above.
* I AGREE TO abide by any conditions attached to the application by the PBPP during the approval process. I understand that failure to comply with these conditions may result in any future applications by me, or my employing or sponsoring organisation, may be refused.
* I CERTIFY THAT all those who have access to health data in this proposal are aware of the requirements of confidentiality and understand that any breach (eg disclosure of confidential information to a person not authorised to receive it) will be reported to the data controller, and in the case of NHS Scotland originated data to Scottish Government eHealth division.
* I CERTIFY THAT that only the persons named in the PBPP form (1.1-1.6) as requiring access to the data will be given access and that the data will not be transferred to anyone else.
* I GUARANTEE THAT no publication will appear in any form in which an individual may be identified without the written permission of that individual, and that I will apply appropriate disclosure control when planning publications involving the data requested.
* I UNDERSTAND THAT the Data Controller, and agents acting on its behalf, reserves the right to inspect the data on the sites where it is being processed.

To be signified by the APPLICANT

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| --- | --- |
| Name (in Capitals): | Date: |

To be signified by the PhD SUPERVISOR (if applicable)

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| --- | --- |
| Name (in Capitals): | Date: |

* I DECLARE THAT (the applicant named above) is a *bona fide* worker engaged in a reputable project and that the data he/she asks for can be entrusted to him/her in the knowledge that he/she will conscientiously discharge his/her obligations, including in regard to confidentiality of the data, as stated in the declaration above.

To be signified by the INFORMATION CUSTODIAN named in Section 1.4 above (where the Information Custodian is not the applicant).

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| --- | --- |
| Name (in Capitals): | Date: |

I ACCEPT the organisation’s obligations and roles with respect to the processing of data for the purposes outlined in this application.

To be signified by the Main Contact for the Lead Organisation named in Section 2.2 above

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| Name (in Capitals): | Date: |