Cover is automatic and a Questionnaire is NOT needed if the research is within the UK & limited to the following activities:

1. Questionnaires, interviews, psychological activity including CBT;
2. Venepuncture (withdrawal of blood);
3. Muscle biopsy;
4. Measurements or monitoring of physiological processes including scanning;
5. Collections of body secretions by non invasive methods;
6. Intake of foods or nutrients or variation of diet (other than administration of drugs).

Refer all other Research involving human participants to the Insurance Officer with the following information to arrange cover (and may incur a charge). Prompt submission of the Questionnaire is recommended, and for an early indication of terms, please submit the research proposal in the absence of full information.

1. Institution: Department:
2. Title of Research:
3. Name(s) of sponsoring organisations:
4. Does the research involve -
	1. investigating or participating in methods of contraception? | Yes/No
	2. assisting with or altering the process of conception? | Yes/No
	3. the use of drugs? | Yes/No
	4. the use of surgery (other than biopsy)? | Yes/No
	5. genetic engineering? | Yes/No
	6. subjects under 5 years of age? (other than activities i-vi above) | Yes/No
	7. subjects known to be pregnant? (other than activities i-vi above) | Yes/No
	8. pharmaceutical product/appliance designed or manufactured by the institution?| Yes/No
	9. work outside of the United Kingdom? | Yes/No

If ‘Yes’ to any of the questions, 4a-i above, and this a follow-on phase, provide details of *SUSAR*s on a separate sheet (fatal or life threatening events)

###

###  Please also provide: the Protocol, the Patient Information & Patient Consent forms, and completed page 2 of this Questionnaire.

### Name: Date:

###

**\***NB:for the purpose of indemnity/cover **Clinical Trial** means**:** an investigation or series of investigations conducted on any person for a **Medicinal Purpose.**

**Medicinal Purpose** means:

1. treating or preventing disease or diagnosing disease or
2. ascertaining the existence degree of or extent of a physiological condition or
3. assisting with or altering in any way the process of conception or
4. investigating or participating in methods of contraception or
5. inducing anaesthesia or
6. otherwise preventing or interfering with the normal operation of a physiological function

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  |   |  |  |  |
| 1. Has NHS Indemnity been provided? |  |  |  | YES | NO |
|  |  |  |  |  |  |
| 2. Will Medical Practitioners be covered by the MDU or other body? | YES | NO |
|  |  |  |  |  |  |
| 3. This section aims to identify those staff involved, their employment contract and the extent of their involvement in the Research. |
|  Name the employer and if an NHS honorary contract is held : |
| (In some cases it may be more appropriate to refer to a group of persons rather than individuals.) |
| Principal Investigator: |  |  |  |  |  |
| Name |  |  | Employer | NHS Honorary contract? Yes/No |
| Activities undertaken:  |  |  |  |
| Others: |  |  |  |  |
| Name |  | Employer | NHS Honorary contract? Yes/No |  |
|  |  |  |  |  |  |
| 4. Please provide any further relevant information here: |  |  |  |
|  |  |  |  |  |  |
| Please copy this form if necessary and continue to list all individuals or groups of staff involved with the Research |