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| **A close-up of a logo  Description automatically generated****Independent Ethics Committee** | Checklist of 21 elements in the informed consent form and written subject information |

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| Checklist of 21 elements in the informed consent form and written subject informationReference: ICH-GCP, Malaysian Guidelines for Good Clinical Practice, 3rd edition, 2011This checklist is a tool to help in determining that the required information is well presented in the Patient Information Sheet and Informed Consent Form. Adherence with this checklist assures compliance with applicable research guideline and regulations. |
|  **Protocol Title:**       |
| **Principal Investigator:**         |

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|  | GCP Informed Consent Elements | Present | Missing | Not applicable |
| 1. | That the study involves research.  |[ ] [ ] [ ]
| 2. | The purpose of the study.  |[ ] [ ] [ ]
| 3. | The study treatment(s) and the probability for random assignment to each treatment.  |[ ] [ ] [ ]
| 4. | The study procedures to be followed, including all invasive procedures.  |[ ] [ ] [ ]
| 5. | The subject’s responsibilities.  |[ ] [ ] [ ]
| 6. | Those aspects of the study that is experimental.  |[ ] [ ] [ ]
| 7. | The reasonable foreseeable risks or inconveniences to the subject and, when applicable, to an embryo, fetus, or nursing infant.  |[ ] [ ] [ ]
| 8. | The reasonably expected benefits. When there is no intended clinical benefit to the subject, the subject should be made aware of this.  |[ ] [ ] [ ]
| 9. | The alternative procedure(s) or course(s) of treatment that may be available to the subject and their important potential benefits and risks.  |[ ] [ ] [ ]
| 10. | The compensation and/or treatment available to the subject, in the event of study-related injury.  |[ ] [ ] [ ]
| 11. | The anticipated prorated payment, if any, to the subject for participating in the study.  |[ ] [ ] [ ]
| 12. | The anticipated expenses, if any, to the subject for participating in the study  |[ ] [ ] [ ]
| 13. | That the subject’s participation in the study is voluntary and that the subject may refuse to participate or withdraw from the study, at any time, without penalty or loss of benefits to which the subject is otherwise entitled.  |[ ] [ ] [ ]
| 14. | That the monitor(s), the auditor(s), the IRB/IEC, and the regulatory authority(ies) will be granted direct access to the subject’s original medical records for verification of clinical study procedures and/or data, without violating the confidentiality of the subject, to the extent permitted by the applicable laws and regulations and that, by signing a written informed consent form, the subject or the subject’s legally acceptable representative is authorizing such access.  |[ ] [ ] [ ]
| 15. | That record identifying the subject will be kept confidential and, to the extent permitted by the applicable laws and/or regulations, will not be made publicly available. If the results of the study are published, the subject’s identity will remain confidential.  |[ ] [ ] [ ]
| 16. | That the subject or the subject’s legally acceptable representative will be informed in a timely manner if information becomes available that may be relevant to the subject’s willingness to continue participation in this study.  |[ ] [ ] [ ]
| 17. | The person(s) to contact for further information regarding the study and the rights of study subjects (e.g. Representative of EthicsCommittee), and whom to contact in the event of the study-related injury (e.g. Investigator).  |[ ] [ ] [ ]
| 18. | The foreseeable circumstances and/or reasons under which the subject’s participation in the study may be terminated.  |[ ] [ ] [ ]
| 19. | The expected duration of the subject’s participation in the study.  |[ ] [ ] [ ]
| 20. | The approximate number of subjects involved in the study.  |[ ] [ ] [ ]
| 21. | The source of the investigational product that may be culturally unacceptable.  |[ ] [ ] [ ]
|      Additional guidance |  |  |  |
| A | If samples of tissue, cells, blood or body fluids are taken, consent must be given to store samples or else you must state that samples will be destroyed at the end of the project. |[ ] [ ] [ ]
| B | Should the subject(s) withdraw, the status of data or biologicalspecimen(s) collected prior to withdrawal should be disclosed to the subject |[ ] [ ] [ ]
| C | The presence or absence of provision for extended access of drugs (for research where a drug(s) is given) |[ ] [ ] [ ]
| D | The language used in PIS and ICF is as non-technical as practical and understandable to the subject or the subject’s legally acceptable representative and the impartial witness, where applicable |[ ] [ ] [ ]
| E | Total amount of blood to be taken for whole should bespecified and expressed in ordinary everyday measure such asnumber of tea-spoon (one tea-spoon is about 5ml) |[ ] [ ] [ ]

Pertinent Format Requirements

**In the Patient Information Sheet, are the following available?**

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|  |  | Present | Missing |  |
| 1 | Title of the research project on every page |[ ] [ ]           |
| 2 | Protocol Number |[ ] [ ]        |
| 3 | Version No. of the PIS on every page |[ ] [ ]        |
| 4 | Sponsor’s Name |[ ] [ ]        |
| 5 | Name of Principal Investigator and Institution |[ ] [ ]        |
| 6 | To mention whether investigational product contain porcine /bovine ingredients |[ ] [ ]        |
| 7 | Page numbers must be expressed on every page |[ ] [ ]        |
| 8 | Date & Time of Signatures |[ ] [ ]        |