**FORM 7.1 E- CHECKLIST FOR ASSESSING THE ADEQUACY OF THE INROMED CONSENT FORM FOR GENETIC STUDIES**

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| CHECKLIST STATEMENT | **YES** | **NO** |
| 1. A statement that the study involves research |  |  |
| 1. An explanation of the purposes of the research |  |  |
| 1. The expected duration of the participant’s participation |  |  |
| 1. A description of the procedures to be followed |  |  |
| 1. A description of any reasonably foreseeable risks or discomforts to the participant |  |  |
| 1. A description of any benefits to the participant and to others that may reasonably be expected from the research |  |  |
| 1. A disclosure of appropriate alternative procedures or courses of treatment, if any, that might be advantageous to the participant |  |  |
| 1. A statement describing the extent, if any, to which confidentiality of records identifying the participant will be maintained |  |  |
| 1. The amount and schedule of all payments |  |  |
| 1. The approximate number of participants involved in the study. |  |  |
| 1. For research involving more than minimal risk an explanation as to whether any compensation is available if injury occurs and, if so, what it consists of, or where further information may be obtained |  |  |
| 1. Anticipated circumstances under which the participant’s participation may be terminated by the investigator without regard to the participant’s consent. |  |  |
| 1. A statement that participation is voluntary and that refusal to participate will involve no penalty or loss of benefits to which the participant is otherwise entitled. A statement that the participant may discontinue participation at any time without penalty or loss of benefits to which the participant is otherwise entitled |  |  |
| 1. A statement that significant new findings developed during the course of the research, which may relate to the participant’s willingness to continue participation will be provided to the participant |  |  |
| 1. An explanation of how to contact the research team for questions, concerns, or complaints about the research. |  |  |
| 1. An explanation of how to contact someone independent of the research team for questions, concerns, or complaints about the research; questions about the participant’s’ rights; to obtain information; or to offer input. |  |  |
| 1. An explanation of whom to contact in the event of a research-related injury to the participant. |  |  |
| 1. A section where the research participants signs to affirm his/ her participants |  |  |
| 1. A section for research team member administering the consent |  |  |
| 1. A section for a witness to sign in case the participant is illiterate. |  |  |