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**Optional Biorepository Consent Reviewer Evaluation Form**

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| **IRB #:** Click or tap here to enter text. | **PI:** Click or tap here to enter text. |
| **Title of Study:** Click or tap here to enter text. |
| **Primary Reviewer:** Click or tap here to enter text. | **Secondary Reviewer:** Click or tap here to enter text. |

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| **GENERAL**  |  | **YES** | **NO** | **N/A** |
| 1. CF is presented in the LSUHSC-NO template or a format that has been approved by the administration of the IRB |  | [ ]  | [ ]  |  |
| 2. Document is written in language understandable to subjects |  | [ ]  | [ ]  |  |
| 3. Length of document is appropriate for the complexity of the study |  | [ ]  | [ ]  |  |
| 4. Text is in a consistent person throughout the consent form |  | [ ]  | [ ]  |  |
| 5. Circumstances of consent process provide prospective participant or lar sufficient opportunity to consider whether or not to participate |  | [ ]  | [ ]  |  |
| 6. Circumstances of consent process minimize the possibility of coercion or undue influence |  | [ ]  | [ ]  |  |
|  | ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **TITLE, CONTACTS, AND SPONSOR** | **YES** | **NO** | **N/A** |
| 1. Study title in consent form matches the title of the protocol |[ ] [ ]   |
| 2. The PI is listed |[ ] [ ]   |
| 3. An emergency contact is provided |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 1: INVITATION TO BE PART OF THE RESEARCH** | **YES** | **NO** | **N/A** |
| 1. Clear reason for why subject is eligible or being asked to participate is present |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 2: WHY IS THE STUDY BEING DONE?**  | **YES** | **NO** | **N/A** |
| 1. Rationale for conducting the study is provided |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 3: WHAT WILL HAPPEN IF I TAKE PART IN THIS STUDY** | **YES** | **NO** | **N/A** |
| 1. Number/amount of blood draws is provided in household terms |[ ] [ ] [ ]
| 2. Description of all procedures is complete |[ ] [ ] [ ]
| 3. Explanation for collection and use of tissues (including DNA analyses) is appropriate |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 4: WHAT SHOULD I KNOW ABOUT GENETIC RESEARCH** | **YES** | **NO** | **N/A** |
| **This section is:** [ ]  **Applicable to this study** [ ]  **Not Applicable to this study (***move to next section***)** |
| 1. Is GINA language included?  |[ ] [ ] [ ]
| 2. If yes, description of how genetic information will be generated/used/analyzed is included |[ ] [ ] [ ]
| 3. If yes, language about whole genome sequencing is included |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 5: WHAT ARE THE RISKS OF TAKING PART IN THIS STUDY?** | **YES** | **NO** | **N/A** |
| 1. All reasonably foreseeable risks are appropriately listed  |[ ] [ ]   |
| 2. Risk of any required routine procedures such as blood draws are listed |[ ] [ ] [ ]
| 3. Statement about unforeseeable risks is included |[ ] [ ] [ ]
| 4. Medical consequences of risks are explained |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 6: ARE THERE ANY BENEFITS TO PARTICIPATING IN THIS RESEARCH?** | **YES** | **NO** | **N/A** |
| 1. Description of potential benefits is appropriate and not overly optimistic |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 7: WHAT OTHER CHOICES DO I HAVE IF I DON’T TAKE PART IN THIS STUDY?** | **YES** | **NO** | **N/A** |
| 1. Subjects are informed that they may choose not to participate |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 8: HOW WILL MY INFORMATION BE KEPT CONFIDENTIAL?** | **YES** | **NO** | **N/A** |
| 1. Subject is made aware that his/her privacy will be protected | [ ]  | [ ]  |  |
| 2. Procedures for privacy protection are described and are adequate | [ ]  | [ ]  |  |
| 3. Time frame of retention of study information/samples is stated | [ ]  | [ ]  |  |
| 4. Agencies/entities/individuals who will have access to participant information and/or study results are identified and are appropriate | [ ]  | [ ]  |  |
| 5. The type of information available to agencies/entities/individuals is described and is appropriate | [ ]  | [ ]  |  |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 9: WILL MY INFORMATION/SPECIMENS BE USED FOR FUTURE RESEARCH?** | **YES** | **NO** | **N/A** |
| 1. Description of how information/specimens will be used in future research is included | [ ]  | [ ]  |  |
| 2. A checkbox and section for initials for future use is included | [ ]  | [ ]  |  |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 10: WILL THERE BE ANY COSTS TO ME FOR TAKING PART IN THIS RESEARCH?**  | **YES** | **NO** | **N/A** |
| 1. Research and/or routine care costs covered by the sponsor is clearly delineated and is appropriate |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 11: WILL I BE PAID FOR TAKING PART IN THIS STUDY?** | **YES** | **NO** | **N/A** |
| 1. Timing and amount of subject payment is indicated and is appropriate |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 12: WHO CAN PROFIT FROM THE STUDY RESULTS?** | **YES** | **NO** | **N/A** |
| 1. Required language related to an investigator COI is provided |[ ] [ ] [ ]
| 2. Language describing potential commercialization of specimens is included |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 13: WHO CAN I CONTACT IF I HAVE QUESTIONS ABOUT THIS STUDY?** | **YES** | **NO** | **N/A** |
| 1. All investigators are listed |[ ] [ ]   |
| 2. A 24-Hour phone number is provided |[ ] [ ]   |
| 3. A research-related injury contact name and number is provided  |[ ] [ ]   |
| 4. A ClinicalTrials.gov statement included |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 14: WHAT IF I NO LONGER WISH TO PARTICIPATE IN THE STUDY?** | **YES** | **NO** | **N/A** |
| 1. Steps to be taken for safe withdrawal from the study is adequately explained |[ ] [ ] [ ]
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 15: YOUR PARTICIPATION IN THIS STUDY IS VOLUNTARY** | **YES** | **NO** | **N/A** |
| 1. Subject’s right to refuse to participate, and with no loss of benefits to which he/she is entitled, is clearly stated |[ ] [ ]   |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |

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| **SECTION 16: YOUR CONSENT** | **YES** | **NO** | **N/A** |
| 1. Statement confirming retention of legal rights after signing the consent is present | [ ]  | [ ]  |  |
| 2. Researchers contact statement is provided | [ ]  | [ ]  |  |
| 3. Chancellor’s contact statement is provided | [ ]  | [ ]  |  |
| 4. All appropriate signature blocks (including date lines) are present | [ ]  | [ ]  |  |
| ***Comments/Deficiencies:*** Click or tap here to enter text. |