**DATABASE/BIOBANK PROTOCOL TEMPLATE**

|  |  |
| --- | --- |
| Title: |  |
| Version Date: |  |
| IRB Number: |  |
|  |  |
| Principal Investigator: |  |
| Sub-Investigator(s): |  |
|  |  |
| Performance Site(s): |  |
|  |  |
| Sponsor: |  |
|  |  |
| Amendment 1 Date: | Amendment 4 Date: |
| Amendment 2 Date: | Amendment 5 Date: |
| Amendment 3 Date: | Amendment 6 Date: |

**STATEMENT OF COMPLIANCE**

This study will be conducted in full accordance with all applicable LSU Health Sciences Center Policies and Procedures and all applicable Federal and state laws and regulations including 45 CFR 46, and the HIPAA Privacy Rule. Any episode of noncompliance will be documented.

The investigators will perform the study in accordance with this protocol, will obtain consent/assent/ HIPAA authorization (unless waivers are granted), and will report unexpected problems in accordance with LSU Health Sciences Center - New Orleans IRB Policies and Procedures and all federal requirements. Collection, recording, and reporting of data will be accurate and will ensure the privacy, health, and welfare of research subjects during and after the study.

**ABBREVIATIONS AND DEFINITIONS OF TERMS**

|  |  |  |
| --- | --- | --- |
|  |  | Insert and delete terms as relevant [Examples Below] |
| EMR |  | Electronic Medical Record (i.e. EPIC) |
| MRN |  | Medical Record Number |
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**PROJECT SUMMARY/ABSTRACT**

**This section should summarize the central elements of the protocol and be able to stand on its own. It should be no more than one page.**

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| --- | --- |
| **Objective(s)** | The purpose of this database/biobank is to…  |
| **Study Population** | *Include a brief description of the population such as health/disease status, gender, age, etc.* |
| **Study Design** | *Present an overview of the study design. For example:* * *Database/ Data Registry or Biobank/Biospecimen Repository*
 |
| **Sample Size** | *Include total number of patients for the study* |

1. **BACKGROUND AND RATIONALE**

This section is based on your research question. The background and rationale should:

* Outline the current knowledge about the research topic
* Include a statement for the need/problem the research topic will address
* Answer why the research needs to be done
* Answer what the relevance is of the research

Include references with citations from the literature. This section should be no more than 2-3 pages.

1. **STUDY OBJECTIVES**

The purpose of this database/biobank is to…

**2.1 Primary Objective (or Aim)**

This should be specific. For example: “The primary object of this study is to develop a biobank of benign tumors for future use.”

**2.2 Secondary Objectives (or Aims)**

The secondary objectives are to [include any additional objectives, if applicable]

1. **INVESTIGATIONAL PLAN**

**3.1 Database/Biobank Design**

Provide a general overview of the database/biobank including the method of collection. The specifics will go in Section 4.

**3.2 Study Sites**

The study will be conducted at [number] sites. Local sites will include…

**3.3 Study Population**

The study team will enroll approximately [number] subjects.

**3.3.1 Inclusion Criteria**

[Examples…

* Had a benign tumor biopsy
* Treated at hospital between January 2014 and December 2023
* Age 18 or older]

**3.3.2 Exclusion Criteria**

[Examples…

* Under age 18
* Tumor biopsy was malignant]
1. **STUDY PROCEDURES**

**4.1 Data/Biospecimen Sources**

This description should be specific but not overly detailed. For example: “EPIC will be queried for demographic information, admission dates, and diagnosis codes. Tissue diagnosis will be obtained from pathology records.”

**4.2 Data Elements/Biospecimen to be Collected**

Provide a listing of all variables that will be collected from records or all sample types that will be collected from the subject.

**4.3 Data Collection and Management**

This section should include the following information:

* How data /samples will be collected, and the system used for storage.
* How confidentiality of the data will be ensures
* Plan for anonymization or de-identification of the data/biospecimen
* Plan for retention and destruction of the data/specimen

**4.4 Sharing of Data or Biospecimen**

Describe the plan for:

* How requests for use of data/biospecimen will be received
* How requests will be reviewed
* If approved, how data/biospecimen will be shared and returned
1. **REGULATORY AND ETHICAL CONSIDERATIONS**

**5.1 Risks to Participant**

Describe the risks of participation in this section. All studies have at least some risk. For chart reviews, the primary risk is the breach of confidentiality of data.

Address how the study design and data management plan will minimize the risks of harm.

**5.2 Potential Benefits to Participant**

Summarize potential benefits of participation, if any.

**5.3 Informed Consent/Assent and HIPAA Authorization**

Either informed consent must be obtained, or a waiver must be requested.

* If consent will be obtained, describe the procedures that will be used to obtain consent (who, when, where, how).
* If a waiver is requested, provide sufficient justification for why the research meets the criteria for a waiver.

**5.4 Safety Reporting**

Describe the plan for reporting any issues to the IRB.

1. **PUBLICATION**

Describe the plan for publication and presentation of the results.

1. **REFERENCES**

[Insert references/citations]