IRB Minimal Risk Study Application

1. Study Title and Investigator Information

Study Title

Principal Investigator Name

Institution/Department

Contact Email

2. Study Summary

Brief Description of the Study

3. Study Purpose and Hypothesis

Purpose of the Study

Hypothesis or Research Questions

4. Study Participants

Description of Participant Population

Inclusion and Exclusion Criteria

Recruitment Methods

5. Study Procedures

Procedures and Data Collection Methods

6. Data Management

How will data be kept confidential?

Data Storage and Security

Duration of Data Retention

7. Risks and Benefits

Describe Potential Risks to Participants

Describe Potential Benefits

8. Informed Consent Process

Describe Process for Obtaining Informed Consent

9. Additional Information

Additional Relevant Information