1. Study Identification

Study Title

Protocol/Study ID

Sponsor (if any)

Principal Investigator

Affiliation

Email

Phone

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2. Study Details

Study Objective

Study Design

Study Location(s)

Sample Size

Study Duration

Study Population

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3. Ethical Considerations

Informed Consent Process

Risks and Benefits

Confidentiality Measures

Involvement of Vulnerable Groups (if any)

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4. Regulatory Approvals

Regulatory Approvals (if any)

Previous Ethics Approval (if any)

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5. Attachments

List of Attachments (e.g., protocol, patient information sheet, consent form, recruitment materials)