

Informed Consent Form for Clinical Trial Participants

Title of Study

Principal Investigator

Institution

Contact Information

Purpose of the Study

Procedures

Risks and Discomforts

Benefits

Confidentiality

Voluntary Participation

Withdrawal

Questions or Concerns

Participant's Statement

- I have read and understood the information above.
- I have had the opportunity to ask questions, and my questions have been answered.
- I consent voluntarily to participate in this study.

Participant Name:

Signature:

Date:

Person Obtaining Consent Name:

Signature:

Date: