

Kelsey-Seybold Clinic Sample Informed Consent to Participate in a Clinical Study

Principal Investigator:
Address

Study Title

Sponsor

Background

Purpose and Design of the Study:

Expected Duration

Description/Procedures

Risks and Discomforts

Information for women of child bearing potential

Benefits of the Study

Alternative methods of Treatment

Subject Rights and Study Withdrawal

Source of Information

Compensation for Injury

Confidentiality

Consent

Name of Subject

Date

Signature of Subject

Date

Name of Witness

Date

Signature of Witness

Date

Signature of Investigator

Date

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Pt. Initials _____