Clinical Research Re-advertising Consent Template

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Study Title:
Principal Investigator:
Purpose of the Study
What does re-advertising involve?
What will I be asked to do?
Risks and Benefits
Confidentiality
Voluntary Participation
Contact Information
Consent
I have read the information above and have had the opportunity to ask questions. I voluntarily consent to be recontacted and/or re-advertised for participation in the clinical research described above.
Participant Name:
Signature:
Date: