Informed Consent Form for Clinical Trials

Title of Study
Principal Investigator(s)
Institution
Purpose of the Study
Procedures
Risks and Discomforts
Benefits
Confidentiality

Voluntary Participation and Withdrawal

Contact Information	
Who to contact for questions about the study:	
Email:	
Phone:	
Participant Statement	
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I have read and understood the information provided above. I have had all my questions answagree to participate in this study.	wered. I voluntarily
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Participant Signature	
Date:	
Investigator Signature Date:	