## **Clinical Trial Informed Consent Form**

Study Title:
Protocol Number:
Principal Investigator:
Site Address:
Phone:
Introduction
Purpose of the Study
Procedures
1. 2. 3.
Risks and Discomforts
•
Benefits
•
Confidentiality
Voluntary Participation and Right to Withdraw
Contact Information
Participant Statement
I have read and understood the information provided above. I have had a chance to ask questions and have received answers to my satisfaction. I voluntarily agree to participate in this study.
Participant Name:

Signature:

Date:			
Investigator Name:			
Signature:			
Date:			